



# STUDIES ON RHIZOSPHERE AND RHIZOPLANE MYCOFLORA OF CHICKPEA

ABSTRACT

THESIS SUBMITTED FOR THE DEGREE OF

**Doctor of Philosophy**

IN

**BOTANY**

BY

**SYED BAQAR IMAM ZAIDI**

T-3613

DEPARTMENT OF BOTANY  
ALIGARH MUSLIM UNIVERSITY  
ALIGARH [INDIA]

1988



# STUDIES ON RHIZOSPHERE AND RHIZOPLANE MYCOFLORA OF CHICKPEA

ABSTRACT

THESIS SUBMITTED FOR THE DEGREE OF

**Doctor of Philosophy**

IN

**BOTANY**

BY

**SYED BAQAR IMAM ZAIDI**

DEPARTMENT OF BOTANY  
ALIGARH MUSLIM UNIVERSITY  
ALIGARH [INDIA]

1988

## ABSTRACT

Studies have been made on the rhizosphere and rhizoplane mycoflora of chickpea (Cicer arietinum L.) inoculated with wilt causing organism Fusarium oxysporum f.sp. ciceri as influenced by certain factors.

There was more fungal population in the rhizosphere of both healthy and wilt infected plants in comparison to non-rhizosphere. There was more population of fungi in the rhizosphere of infected chickpea plants with higher frequency values and R:S ratio than healthy counterparts. It is likely that the roots of infected plants have higher concentration of total free amino acids and sugars than healthy ones which might be favouring activity of fungi in infected rhizosphere. The roots of infected plants also have higher concentration of phenols and O-dihydroxy phenols.

Different cultivars of chickpea differed in the rhizosphere mycoflora both qualitatively and quantitatively. There were some species of fungi found in the rhizosphere of certain cultivars only. Differences were also observed with respect to free amino acids, phenols, O-dihydroxy phenols and sugars in the root extracts from different cultivars. The cultivars

which harboured more fungi and have higher frequency of F. oxysporum f.sp. ciceri in the rhizosphere and rhizoplane have usually low concentration of free amino acids, phenols and O-dihydroxy phenols but higher concentration of sugars. The reverse was true with cultivars which harboured lower number of fungi together with lower frequency of F. oxysporum f.sp. ciceri. The number of amino acids have been relatively high in the cultivars susceptible to wilt causing organism i.e. JG-62 and H-208 as compared to BG-212 and JG-74 which are less damaged. Qualitatively, Asparagine and Histidine which have been detected in the roots of JG-62, H-208, BG-309 and JG-315 are known for their stimulatory effect on the growth of fungi while cysteine in the BG-212 and JG-74 might be contributing towards the resistance of plants against the disease. By and large, more fungi were detected in H-208, JG-62 and BG-309 than JG-74, BG-212 and JG-315. Moreover, the frequency of pathogenic fungi including F. oxysporum f.sp. ciceri was more in the rhizosphere and rhizoplane of JG-62 and H-208 and that of antagonistic fungi particularly T. viridae in BG-212 and JG-74. The fungus population was also high in the rhizosphere of H-208 and least in JG-74 in both inoculated and uninoculated plants.

There was an increase in population of fungi, R:S ratio and number of fungi in the rhizosphere and rhizoplane with age



of both inoculated and uninoculated plants. The highest rhizosphere activity was observed at flowering and fruiting stage followed by a decline at senescence. Different plant growth stages also influenced the species composition of rhizosphere mycoflora. However, Aspergilli and other saprophytic fungi constituted dominant flora throughout the growth period in uninoculated plants. On the other hand, F. oxysporum f.sp. ciceri and other parasitic fungi dominated in the rhizosphere of inoculated plants. The concentration of total free amino acids, phenols, O-dihydroxy phenols and sugars also increased with increase in age of plant being maximum in 75 to 90 days old plants followed by a decline at senescence.

Foliar spray with different growth promoting substances by and large resulted in an increase in the population and frequency of mycoflora however with maleic hydrazide, there has been a decrease in the majority of fungi including wilt pathogen F. oxysporum f.sp. ciceri except A. fumigatus, A. niger, Trichoderma lignorum. Foliar sprays with urea exhibited stimulatory effect on the activity of saprophytic as well as parasitic fungi in the rhizosphere and rhizoplane of both inoculated and uninoculated plants but with muriate of potash most of the saprophytic fungi increased while that of parasitic fungi decreased. Sprays with different fungicides brought about an inhibition in the population of rhizosphere fungi. In addition to above, the frequency of majority

of the fungi including F. oxysporum f.sp. ciceri also decreased due to foliar sprays with fungicides but that of saprophytic fungi such as Aspergillus clavatus, A. flavus, A. fumigatus and T. viridae increased. On the other hand, spray with streptomycin generally stimulated the rhizosphere fungal flora.

Incorporation of different fertilizers in the soil resulted in an increase in the rhizosphere population of fungi in uninoculated plants. Similarly, soil amendment with oil cakes, by and large, increased the rhizosphere and rhizoplane mycoflora except mahua cake. But in inoculated ones, there was a decrease in the rhizosphere and rhizoplane mycoflora activity in all the treatments except with urea and superphosphate. Different fungicides when incorporation in soil brought about a decrease in rhizosphere population of fungi both in inoculated and uninoculated plants. The frequency of most of the saprophytic and parasitic fungi including F. oxysporum f.sp. ciceri increased with urea and superphosphate. However, with muriate of potash and different oil cakes, the frequency of most of the saprophytic fungi increased and that of parasitic fungi including F. oxysporum f.sp. ciceri decreased. With fungicides, the frequency of majority of the fungi in the rhizosphere and rhizoplane decreased. F. oxysporum f.sp. ciceri was either not detected or its frequency was low in soil treatment with castor cake, mahua cake, neem cake, bavistin, benlate and vitavax.



# STUDIES ON RHIZOSPHERE AND RHIZOPLANE MYCOFLORA OF CHICKPEA

THESIS SUBMITTED FOR THE DEGREE OF

**Doctor of Philosophy**

IN

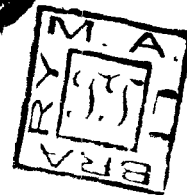
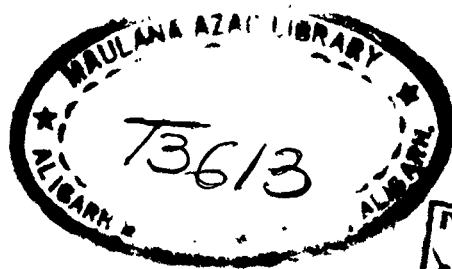
**BOTANY**

BY

**SYED BAQAR IMAM ZAIDI**

DEPARTMENT OF BOTANY  
ALIGARH MUSLIM UNIVERSITY  
ALIGARH [INDIA]

1988



- 1 JUN 1989

THESIS SECTION



T3613

CHECKED 1996-97

Dedicated to the  
Memory of  
Dr. Dhirendra Prakash

*S. K. Saxena*  
M.Sc., Ph.D., F.B.S., F.P.S.I.  
Professor



Telephone : { Office : 4984  
                  { Res. : 3161

DEPARTMENT OF BOTANY  
ALIGARH MUSLIM UNIVERSITY  
ALIGARH-202001 (INDIA)

Dated....20.6.1988.....

CERTIFICATE

This is to certify that Mr. Syed Baqar Imam Zaidi has worked in this department as a research scholar under my supervision and guidance. His work "Studies on rhizosphere and rhizoplane mycoflora of chickpea" is upto date and original. He is allowed to submit his thesis for the award of the degree of Doctor of Philosophy in Botany.

*S. K. Saxena*  
( S. K. ~~SAXENA~~ )

## C O N T E N T S

### PAGE NO.

#### ACKNOWLEDGEMENTS

CHAPTER	I :	Introduction and Review of Literature	...	1
	II :	Materials and Methods	...	37
	III :	Experimental Results	...	51
	IV :	Discussion	...	100
		SUMMARY	...	119
		REFERENCES	...	123

-----

## ACKNOWLEDGEMENTS

It is with a deep sense of gratitude that I place on record my indebtedness to **Dr. Dharendra Prakash** (Late), Reader, Department of Botany, Aligarh Muslim University, for proposing the problem on which this work was carried out. The problem proved to be of vital interest. His able guidance during his life time was great source of inspiration to me. Though he has passed away, I shall always remember his kind hearted treatment, able guidance and unceasing encouragement. He and his virtues will forever be a guiding source in my life.


I am deeply indebted to **Professor S.K. Saxena**, Department of Botany, Aligarh Muslim University, Aligarh who very kindly stepped in to guide me after the sad demise of Dr. Prakash. His valuable supervision, sympathetic attitude, constructive suggestions and his taking consistent pains towards the preparation of this thesis are a cornerstone of the present work.

I am equally thankful to **Professor Khalid Mahmood**, Chairman Department of Botany for his help and for providing the necessary facilities.

I want to extend my grateful thanks to **Mr. S. Zakiul Hasan Zaidi** and **Dr. S.Q.A. Naqvi** of the Department for taking interest in the problem at all times and also **Dr. V.S. Verma**, Division of Mycology & Plant Pathology, Indian Agricultural Research Institute (I.A.R.I), New Delhi for helping me on several occasions.

Thanks are also due to all of my **friends** and **colleagues** for their many fold help, cooperation and critical suggestions during the course of study.

The award of Senior Research Fellowship by C.S.I.R., New Delhi is also gratefully acknowledged.



(Syed Baqar Imam Zaidi)



# **Chapter One**

## **Introduction and Review of Literature**

## CHAPTER - I

### INTRODUCTION AND REVIEW OF LITERATURE

1.1 Rhizosphere has been defined by Hiltner (1904) as that region of the soil which is subject to the influence of plant roots and is characterized by greater microbiological activity than the soil away from the plant roots. Later terms like "Edophosphere" for the rhizosphere of Hiltner and "Histosphere" for the root surface microflora have been coined (Perotti, 1926). Clark (1949), however, suggested "Rhizoplane" for the ecological niche or habitate provided for microorganisms by root surface. However, this field of study remained somehow neglected until Starkey (1929) exhibited a relationship between the soil microorganisms and the green plants. Since then considerable work has been done on the rhizosphere microflora of various economically important plants such as chickpea (Sondhi and Sinha, 1963; Gujrati, 1969; Mathur and Chauhan, 1972; Khan and Prakash, 1982; Satyaprasad, 1982 and Satyaprasad and Ramarao, 1983), pigeon pea (Agnihothrudu, 1955 and 1957; Rai and Upadhyay, 1980 and 1986 and Singh and Bhargava, 1981), soyabean (Rouatt et al; 1963; Mohamed, 1985; and Dutta and Deb, 1986), pea (Timonin, 1940; Stenton, 1958; Luke and Devi, 1975 and Son et al., 1985), bean (Ito and

U1, 1975 and Devet et al., 1980), broad bean (Youssef and Mankarios, 1974), cluster bean (Bahadur and Sinha, 1965), lentil (Gujrati, 1969), groundnut (Krassilnikov et al., 1933; Gangawane and Deshpande, 1975 and 1977 and Gunasekar and Rao, 1982), alfalfa (Thom and Humfeld, 1932; Timonin, 1940; Ishizawa et al., 1957; Stelfox and Williams, 1980 and El-Hamolawi and Erwin, 1986), cowpea (Murthy and Raghu, 1976), wheat (Simmonds and Ledingham, 1937; Timonin, 1940; Warcup, 1957; Herr, 1957; Peterson, 1958; Moskovets, 1957; Neal et al., 1970; Chrzanowski, 1976; Hornby and Brown, 1977; Smiley, 1978; Neate et al., 1981; Ashraf, 1981; Mohamed, 1985 and Ermekova and Abirova, 1985), barley (Ishizawa et al., 1957; Kirilenko, 1970 and 1973; Murthy and Raghu, 1976; Loper et al., 1984; Gerhordson et al., 1985 and Ansari, 1982), maize (Moskovets and Zhdanova, 1960; Kulshrestha, 1969; Kulshrestha et al., 1977 and Annapurna and Rao, 1983), pearl millet (Natrajan, 1972 and Ashraf, 1981), oat (Timonin, 1940; Warcup, 1957; Chester and Parkinson, 1959; Kirilenko, 1970 and 1973; Karimbaeva and Sizova, 1976; Gerhardson et al., 1985), rye (Thom and Humfeld, 1932), cotton (Clark and Thom, 1939; Patel and Iyer, 1961; Harrison and Beckman, 1982, and Moubasher and Abdel Hafeez, 1986), flax (Berezova, 1941 and Timonin, 1940b), strawberry (Katznelson and Richardson, 1948), potato (Atkinson and Robinson, 1955; Loper et al., 1984 and Choroszewski, 1985), timothy (Ishizawa, 1957), clover (Lime, 1984 and Wong et al.,

1986), redclover (Peterson, 1958), birch (Karimbaeva and Sizova, 1976), yellow birch (Ivarson and Katznelson 1960), lupin (Krivets, 1975), yellow lupin (Rataj-Guranowsky, 1981), onion (Parkinson and Clark, 1961; Fenwick, 1973 and Ashour et al., 1980), sesame (Monoharacharya et al., 1977) sunflower (Monoharacharya et al., 1977 and El-Hissy et al., 1980), ornamental plants (Yuen and Schroth, 1986), Medicago sp. (Bretage and Kollmorgen, 1986), cauliflower (Rao and Sharma, 1976), pine (Karimbaeva and Sizova, 1976 and Peno et al., 1977), spruce (Karimbaeva and Snova, 1976 and Stefurak, 1976), garlic (Bertoldi et al., 1978), Citrus sp. (Ali et al., 1979), pomegranate (Ali et al., 1979), grape vine (Ali et al., 1979), Dolichos biflavus (Behera and Patnaik, 1980), tomato (Dwivedi and Pathak, 1981; Chandra et al., 1982 and Loper et al., 1984), Cynodon dactylon (Wadhawani and Mehrotra, 1982), coffee (Venkatasubbaiah et al., 1984), date palm (Ali et al., 1979), Chrysanthemum (El-Hissy et al., 1980), Nigella sativa (El-Hissy et al., 1980), Dhatura inoxia (El-Hissy et al., 1980), Hyosymus muticus (El-Hissy et al., 1980), Cycas revoluta (Wadhwani and Srivastava, 1985), radish (Loper et al., 1984; Gerhardson et al., 1985), sugarbeet (Loper et al., 1984; Lee et al., 1985 and Vesely, 1985 and 1986), Doctylus glomerata (Kutrzeba, 1984), lettuce (Gerhardson et al., 1985), rape (Gerhardson et al., 1985), banana (Goos and Timonin, 1962), cucumber (Vancura and Hovadik, 1965 and Hong, 1969) sugarcane (Prakash, 1967;

Robinson, 1970, Kamal and Singh, 1974; Kishan et al., 1982 and Kao and Hsich, 1985), tobacco (Luke and Devi, 1972) and Abelmoscus esculentus (Dayal and Srivastava, 1975 and Srivastava and Dayal, 1986). These studies have indicated an interaction between soil microbes, soil borne pathogens and higher plants (Starkey, 1929; Timonin, 1940; Agnihothrudu, et al., 1955; Agnihothrudu, 1957, 1958 and 1960; Skilyagina, 1967; Mukerji, 1968; David and Baker, 1970; Linderman, 1970; Reddi and Rao, 1972; Mall, 1973; Raicu and Stan, 1975; Wnekowski, 1975; Newman, 1978; Arora and Dwivedi, 1979; Singh et al., 1979; Marois and Mitchell<sup>1</sup>, 1981; Stenek, 1981; Sheata et al., 1983; Kurtzeba, 1983; Upadhyay and Bharat Rai, 1983; Gokulapalam, 1984; Bristo and Wyllie, 1984; Mohamed, 1985; Beggle-Ristaino and Papavizas, 1985; Kao and Hsich, 1985; Zozzerini and Tosi, 1985; Camporota, 1985; Gerhardson et al., 1985; Vesely, 1986; Upadhyay and Rai, 1986; Yuen and Schroth, 1986; Lockwood, 1986; Arora, 1986; Srivastava and Dayal, 1986 and Phillips, 1986). It has now been shown that rhizosphere microflora have a definite role to play in the development of root diseases (Agnihothrudu, 1959, Timonin, 1966; Srivastava, 1968; Vishwanath et al., 1969; Garrat, 1970; Voronkevich et al., 1972; Louvet, 1972; Samiilenko and Gvozdyak, 1972, Goel and Mehrotra, 1974; Berestetskii, 1974; Pieczarka and Lorbeer, 1975; Zhukovskaya, 1976; Gilligan, 1979; Patel et al., 1981; Rotaj-Guranowsky, 1981; Furuya, 1981; Kutrzeba, 1984; Lim and

Cole, 1984; Choroszewski, 1985; Bretag, 1985; Gagne et al., 1985; Mohamed, 1985; Kao and Hsich, 1985; Vesely, 1985 and 1986; Upadhyay and Rai, 1986; Wong et al., 1986; Sakthival et al., 1986; Agarwal, 1986 and Ermekova and Abirova, 1986). The literature on various aspects of rhizosphere mycoflora has been reviewed by Smith (1948), Katznelson et al. (1948), Clark (1949), Lochhead (1952), Starkey (1958), Katznelson (1961 and 1965), Rovira (1965), Parkinson (1967) and Srivastava (1973 and 1974).

The rhizosphere effect infact becomes evident right from the germination of seed and emergence of roots in the early stages of plant growth. Timonin (1940) observed the development of rhizosphere microflora within three days of seed germination, but further development depended upon the normal growth of the plant. Starkey (1931), while studying the rhizosphere microorganisms of several agriculturally important plants, observed<sup>a</sup> higher number of microorganisms in soil in the immediate vicinity of the roots than in soil away from the roots. Thom and Humfeld (1932) noticed 40-70 times more fungi in the rhizosphere of corn plants than in the non-rhizosphere. Similar observations have been made by Ishizawa et al. (1957), Reddy (1959), Maliszeska and Moreau (1959), Ivarson and Katznelson (1960), Strzekzyk (1961), Rouatt and Katznelson (1961), Zagallo and Bollen (1962), Prakash (1967), Kulsherestha

(1969), Luke and Devi (1975), Dayal and Srivastava (1975), Prakash et al. (1979), Ashraf (1981) and Ansari (1982).  
<sup>A higher</sup> Higher percentage of <sup>(inhibitory to —???)</sup> ~~inhibitory~~ bacteria <sup>the</sup> from rhizosphere of leguminous plants (Gagne et al., 1985) and a low population<sup>s</sup> of fungi in the rhizosphere<sup>s</sup> of Rauwolfia serpentina (Padma and Mukerji, 1972), Allium sativum L. (Yasmeen et al., 1982) and Leucaena leucocephala (Megharaj et al., 1987) have been attributed to ~~(be due to)~~ toxic principle<sup>s</sup> present in root exudates.

Cataska et al. (1960), Dickinson and Paugh (1965a,b), Hongyal-Balul (1983) and Singh et al. (1984) pointed out that there was a relation between the colonization of roots and seed mycoflora. On the other hand, no such relation was observed by Peterson (1959). He, however, concluded that for colonization of roots soil is the primary source of inoculum. Rhizosphere<sup>s</sup> <sup>have</sup> has been shown to provide stimulatory effect<sup>s</sup> to certain fungi (eg. Melanospora breviostrata Moron, Aspergillus gignentius Wehmer and Odecephalum corophilum Kobayasi (Agnihothrudu et al., 1955), Rhizopus orrhizus Fischer, R. nodosus Namy, Choanephora cucurbitarum (Berk and Ray Thaxt and 16 ascomycetes and 50 imperfect fungi excluding <sup>ium spp.</sup> Penicillia (Agnihothrudu, 1957, 1958 and 1960), on Fusarium oxysporum Schl. (Wnekowski, 1975), on F. oxysporum f. sp. vasinfectum (Steve and Curl, 1980) on saprophytic colonization of F. udum (Upadhyay and Bharat Rai, 1983)). The rhizosphere

? clarify preposition? What? as Fusarium??

effect has, however, been found to persist even after 30 days of the harvest of crop (Majumdar and Bhide, 1970). The rhizosphere microflora <sup>are</sup> ~~is~~ influenced by several factors, such as plant growth stages (Krossilnikov, et al., 1936; Riviera, 1959; Gujrati, 1969; Dayal and Srivastava, 1975; Sudha Mall, 1977; Odunfa and Oso, 1979; Khan and Prakash, 1982; Satyaparasad 1982 and Mohamed, 1985); plant type (Tolle and Rippel-Badles, 1957; Bagyaraj and Rangaswami, 1966; Oblisami et al., 1971; Karimbaeva and Sizova, 1976; Rataj-Guranowsky, 1981; Parker, 1985 and Gangawane, 1985); root characters (Timonin and Lochhead, 1948; Sivasithampam and Paker, 1979 and Wadhwani and Srivastava, 1985); nature of root exudates (Rovira, 1962; 1965 and 1969 and Gussin and Lynch, 1983); environmental factors (i.e., type, reaction and fertility of the soil, soil profiles and organic and inorganic content of soil) (Peterson, 1958; Joshi, 1982; Vardavakis, 1986 and Dick and Ali-shatayah, 1986); percentage of moisture content (Clark, 1940; Venkatesan, 1962; Gujrati, 1968 and Bissett and Parkinson, 1979); <sup>light</sup> light (Roviera, 1959; Srivastava, 1971); <sup>temperature</sup> temperature (Rouatt et al., 1963; Namordoze, 1975 and Krikun, 1985); and seasonal variation (Katznelson, 1946; Millar and Boothroyd, 1962; Marks et al., 1975; Singh and Bhargawa, 1981 and Vardavakis, 1986); soil amendment with fertilizers, nematicides, pesticides and fungicides (Clark and Thom, 1939; Absalyamova, 1963; Mishra, 1972; Dath, 1982; Singh and Prasad, 1986 and Yarden, 1986);



foliar spray with chemicals (Rovira, 1959; Balasabramanian and Rangaswami, 1973 and 1975 and Bagyaraj and Rangaswami, 1982); and interactions involving soil microbes and soil borne plant pathogens of higher plants (Skilyagina, 1967; Wnekowski, 1975; Ellingboe, 1983 and Yuen, 1986). <sup>These actions have been</sup> and it ~~has~~ been modified to have favourable effect<sup>s</sup> on ~~the~~ crop<sup>s</sup>.

## 1.2 FACTORS AFFECTING RHIZOSPHERE

*↑ add references to document this point.*

### 1.2.1 Plant Growth Stages

Age of the plant profoundly influenced saprophytic and parasitic rhizosphere mycoflora qualitatively as well as quantitatively (Vesely, 1985). By and large, the rhizosphere activity increased with the increase in the age of plant, attaining highest activity at the peak of vegetative growth of the plant (Timonin, 1940; Khasanov, 1967 and Sudha Mall, 1977). Krossilnikov et al. (1936) observed<sup>a</sup> rise in the population of microorganisms<sup>s</sup> during the early growth period and at the time of formation of fruit as well. Rao (1962) also found two peaks in the increase in the rhizosphere (ie. at the time of flowering and at senescence). Gujrati (1969) and Khan and Prakash (1982) reported higher rhizosphere fungal population at flowering and fruiting stages followed by a decrease, but there was again an increase at senescence stage of lentil and gram. Luke and Devi (1975) observed an increase in rhizosphere activity for 30 days of vegetative growth and

during flowering and fruiting of pea plant.<sup>S</sup> Riviera (1959) noticed that the rhizosphere activity of wheat reached a distinctive peak at the tillering stage influencing fungi more than bacteria. However, Khasanove (1967), Dayal and Srivastava (1975) and Behera and Patnaik (1981) observed that the population of fungi in the rhizosphere of different plants increased upto <sup>the</sup> flowering stage followed by <sup>a</sup> reduction as the plant attained senescence. Two peaks in the rhizosphere activity during the growth of <sup>the</sup> plant <sup>were</sup> also observed by Odunfa and Oso (1979). Rhizopus, Pythium and other phycomycetes fungi dominated at the seedling stages of growth but were succeeded by Fusarium, Curvularia, Phoma and many sterile mycelia at a later stage. Mohamed (1985) observed that <sup>the</sup> population<sup>S</sup> of Streptomyces sp. in the rhizosphere<sup>S</sup> of wheat and soyabean <sup>were</sup> ~~was~~ scarce during the week of seed germination, gradually increasing to maximum at 3-4 wks and then declining. Mishra and Kamal (1972) recorded highest population<sup>S</sup> of fungi at maturity in the rhizosphere<sup>S</sup> in Euphorbia hirta, E. thymifolia, Phyllanthus sp., P. niruri and Smilax.<sup>sp</sup> Satyaprasad (1982) (although observed an increase in the rhizosphere effect in both resistant and susceptible chickpea varieties with increasing age but the population was <sup>greater</sup> ~~more~~ in susceptible varieties.

### 1.2.2 Root Character

Dadalaury and Kotishvili (1975) concluded that the distribution of fungi depended upon the characters of root,<sup>S</sup>

at <sup>s</sup> particular age<sup>s</sup>, and <sup>that</sup> not all the parts of the plant are identical in supporting the rhizosphere population. Timonin and Lochhead (1948) reported more rhizosphere activity in the central or crown portion of the root, which decreased with increasing distance both in horizontal and in vertical direction from the base of the stem. Both qualitative and quantitative differences in the rhizosphere population were observed at different depths of soil profile (Bagyaraj and Rangaswami, 1966; Hussey, 1977; Behera and Mukerji, 1979 and Choe et al., 1986). Sivasithampam and Paker (1979) observed higher population of bacteria, actinomycetes and fungi in the rhizosphere of seminal roots of wheat than the nodal roots. Differences in the rhizosphere activity were also observed between corolloid and normal roots of Cycas revoluta Thomb (Wadhwani and Srivastava, 1985).

### 1.2.3 Plant Type

Rhizosphere<sup>s</sup> of different plants have selective effects on fungi. Certain fungi from the rhizosphere of wheat failed to develop in the rhizosphere of other plants (Tolle and Rippel-Badles, 1958, 1975). The rhizosphere of lucerne differed from that of cotton and that of spring wheat from winter wheat (Moskovets, 1957). Bagyaraj and Rangaswami (1966); Youssef and Mankarous (1968); Oblisam et al. (1971); Srivastava and Mishra (1971); Karimbaeva and Sizova (1976);

can't find reference omitted? Aick to "References" Incorrect citations!!

✓ n.g.s. can not find reference omitted?

Not cit. in refer- and p. Adh.

Rataj-Guranowsky (1981), Parker (1985) and Gangawane (1985) concluded that fungal species composition in the rhizosphere depends upon the type of the plant. Even differences in the rhizosphere microflora were observed between varieties of grapevine (Isarlishvili, 1957). Gujrati (1969) attributed the differences in the rhizosphere microflora of lentil and Cicer arietinum L. to be due to differences in the amino acid contents of root exudates of the two plants. Different rhizosphere mycoflora under different crop covers were also reported by Tandon and Tiwari (1982). Mohamed (1985) found larger number of antagonistic Streptomyces sp. in wheat roots than in soyabeans.<sup>roots</sup>

#### 1.2.4 Root Exudates

It is true that root exudates play a definite role by exuding substances from roots. The root exudates therefore, are found to differ from plant to plant and from young to old plant. There has been immense amount of literature which has been reviewed by Rovira (1962, 1965 and 1969), Scroth and Hildebrand (1964) and Wood (1960). There has been much work on the nature and magnitude of root exudates, the dimension of the rhizosphere, and the kind, sequences and extent of colonization of root surfaces by microbes (Parkinson, 1967 and Bowen, 1979). Knudson (1920) demonstrated that peas and maize plants grown under aseptic conditions in<sup>a</sup> medium containing

sucrose produce considerable quantity<sup>ies</sup> of reducing sugars. Organic nitrogen has been shown to be exuded from roots of maize growing under sterile conditions (Lyon and Wilson, 1921). Various kinds of compounds such as vitamins (thiomine and biotin) from flax seedlings (West, 1939); glucose, flavonose and nucleotides from peas (Lundegradh and Stenlid, 1944); nucleic acids from germinating pea seeds (Fries and Forsman, 1951); amino acids and a reducing sugar similar to ~~that of~~ glucose from roots of tomato, soyabean, barley and oat (Katznelson, 1954); amino acids from pea roots (Rovira, 1956a); glucose, fructose, ribose, galactose and xylose from clover roots (Smith, 1966); lucine, <sup>space</sup>methionine, glutamic acid, aspartic acid, cystine, phenylalanine, arginine (amino acids); rhamnose, arabinose, glucose, fructose, sucrose and raffinose (sugars) from roots of some leguminous crops (Roy and Dwivedi, 1967); organic acids and sugars from the root exudates of Sorghum vulgare and Brassica juncea (Bhuvaneshwari and Subba Rao, 1957); fatty acids (Palmaic, stearic and oleic acid) from bean (Papavizas and Kovacs, 1972) have been reported. Qualitative and quantitative changes in the nature of root exudates have also been found between resistant and susceptible varieties of Hibiscus esculentus (Afifi, 1976). Similar, results have been reported by Furgal-Wegrzycka (1984), Cahill et al. (1985), Mikzak et al. (1985a) and Chaudhary et al. (1986). Satyaprasad and Rama Rao (1983) found qualitatively more amino acids in

the exudates of susceptible in comparison to resistant variety of chickpea. Aspartic acid, asparagine and histidine were detected only in the exudates of the susceptible variety while cystine and galactose were consistently present in the exudates of the resistant variety. Mikzak et al. (1985b), found that aspartic acid was higher in susceptible cultivars while methionine, glycine, leucine and proline <sup>were higher</sup> in resistant cultivars of hops.

Srivastava (1973) demonstrated a direct relationship between rhizosphere microflora and amino acids in the root exudates and root extracts of Echinochloa curs-galli and Papsalum seorbiculatum. Sullia (1973), while studying the effect of root exudates and root extracts of Cassia tora and Crotolaria medica, observed that they stimulated the growth of most of fungi except Trichoderma lignorum (Tode) Harz. Buxton (1957) observed that oospores of Pythium mamillatum Merris. failed to germinate in non-rhizosphere soil or in distilled water but germinated in the presence of growing turnip seedlings. Similarly Coley-Smith and Hickman (1957) observed that the sclerotia of Sclerotium cepivorum Berk. germinated in the presence of onion roots. Roy and Dwivedi (1967) found complete inhibition of conidial germination of Helminthosporium sativum Pammel, King and Bakke and Fusarium culmorum (Smith) Sacc. on glass slides in sterilized soil, but <sup>the</sup> majority of conidia germinated in the vicinity of wheat seedlings. Youssef

very interesting  
not studied in  
Buxton 1957  
mixed up  
with mine  
by Buxton  
(Nature Land)  
190:613-614.

and Mankarios (1969) reported that root exudates of broad bean and cotton stimulated spore germination and growth of rhizosphere fungi. Sharma and Sinha (1974) observed stimulation in germination of fungal spores in the exudates of linseed roots. Schench and Stotzky (1975) pointed out that volatile compounds from germinating seeds of six plant spp. increased growth of several fungi and bacteria, including Erwinia carotovora (var. carotovora), Agrobacterium tumefaciens, Fusarium oxysporum f. sp. conglutinans and Trichoderma viride. Karimbaeva and Sizova (1977) observed a positive relationship of concentration of root exudates on growth of fungi. Gussin and Lynch (1983) found that Fusarium culmorum caused more inhibition of plant growth of wheat, barley and rye grass when grown on root exudates than when present as a spore suspension in sand. El-Hamalawi and Erwin (1986) found <sup>a three to four</sup> (3-4) fold higher spore germination of Phytophthora megasperma f. sp. medicaginis in root extracts and root exudates of alfalfa than in water.

*not in water*  
Tu (1972), while studying (on) seed and root exudation in relation to Rhizoctonia damping off of various crops, concluded that the incidence of disease appeared to be mainly affected by the nature of the exudates. Afifi (1976) observed a correlation between the pathogenic potentialities of Fusarium oxysporum, F. moniliforme (Gibberella fujikuroi), F. solani and F. semitectum and volatile and gaseous exudates of germinating

seeds or roots of Hibiscus esculentus; ~~and found that~~ exudates from <sup>the</sup> susceptible cultivar stimulated conidial germination of all ~~the~~ four fungi whereas those from the resistant cultivar stimulated germination of <sup>only</sup> F. moniliforme. Satyaprasad and Rama Rao (1983) observed that root exudates from a susceptible Cicer arietinum L. cultivar stimulated mycelial growth and germination of conidia and chlamydospore of <sup>the</sup> wilt pathogen (Fusarium oxysporum f. sp. ciceri), while those from a resistant cultivar exhibited inhibition. Furgal-Wegrzycka (1984) found that mycelial growth and sporulation of Fusarium spp. and Ascochyta spp. were inhibited by extracts of resistant but not those of susceptible pea lines.

Burden et al. (1974) attributed natural resistance in pea and bean against root rot to be due to antifungal compounds secreted by roots. Similarly, Salt (1982) observed <sup>plant</sup> wilt resistance in fababean was related to  $\beta$ -alanine content of seed, root and root exudates. Mikzak et al. (1985a) reported that contents of phenols, chlorogenic acid and monophenol/polyphenol ratio were indicators of resistance to Verticillium albo-atrum and Fusarium sambucianum (Gibberella pulicaris). The concentration being higher in resistant hops. Mikzak et al. (1985b) also found that in hop cultivars susceptible to Verticillium albo-atrum and Fusarium sambucianum (Gibberella pulicaris), the level of aspartic acid was higher (but <sup>?</sup> than? that of glycine,



leucine, and proline in the resistant cultivars. Deacon and Mitchell (1985) attributed resistance of oat roots to Allomyces arbuscula, Aphanomyces sp., Phytophthora cinnamomi, Pythium aphanidermatum, P. arrhenomanes, P. graminicola, P. intermedium, P. ultimum var. sporangiferum and Spórolegnia litoralis to be due to saponin in the root.

#### 1.2.5 Soil Type

Soil plays an important role for growth and metabolism of <sup>plants, and</sup> plant, thereby influencing <sup>PS</sup> indirectly the rhizosphere microflora (Dick and Ali-Shtaych, 1986). Peterson (1958), while comparing soil fungal flora around wheat and red clover, concluded that there was no difference in <sup>the write out</sup> (R:S) ratio of plate counts of red clover in the three soils, <sup>e</sup> however, for wheat <sup>the</sup> R:S ratio was 34 in the acidic sandy loam and of 10 in neutral clay loam. <sup>A</sup> Fusarium sp. was found on red clover on the root surface in acid soil, whereas <sup>a</sup> Cylindrocarpus sp. <sup>was found</sup> in the acidic and neutral soils. <sup>A</sup> Gliocladium sp. was not isolated from roots of plants growing in acid or alkaline soil but was present in neutral soil. With wheat grown in acid soil, Fusarium sp. <sup>?</sup> again predominated, but in the alkaline soil <sup>a</sup> Gliocladium sp. <sup>?</sup> Rapidly <sup>sporulating</sup> types such as Penicillium sp. were most numerous in both acid and neutral rhizosphere <sup>s</sup> and root free soils for both plants, but not in chalky soil. This kind of preferential selection of fungi in different soil types has been observed

by Welte and Trolldenier, 1961; Zhdanova, 1963; Joffe, 1969; Downes, 1972; Zgorkiewicz and Mackiewicz, 1975; Bisset and Parkinson, 1979; Behera and Mukherji, 1979; Joshi, 1982, Vardavakis, 1986; Dick and Ali-Shtayah, 1986. However, Kamal and Singh (1969) found no significant effect of soil factor on distribution of fungal population.

#### 1.2.6 Soil Moisture

Early studies by Clark (1940) and Timonin (1940) showed increased microbial counts in the rhizosphere of wheat and flax, respectively, as the soil moisture content decreased. Similar observations were made later by Clark (1948) with soybean. Tresner (1954) observed that the size of the soil microfloral populations was correlated with the moisture and organic matter content of the soil. Venkatesan (1962) obtained highest counts of bacteria, fungi and actinomycetes in the rhizosphere of rice in soil with 20% and 40% moisture, but the R:S ratio increased with greater moisture content until saturation, the high R:S values under submerged conditions were probably due to low soil counts. Sondhi (1964) observed a greater fungal species and higher population in the rhizosphere of Cicer arietinum at 15% moisture than at 25% or 35%. Soil analysis of Cicer arietinum and Lens culinaris by Gujrati (1968) showed that a change in the moisture content

had a pronounced effect on the <sup>f</sup>ungal flora. Similar results were obtained by Bissett and Parkinson, 1979 and Manoharacharya and Bilolikar, 1979.

### 1.2.7 Light and Temperature

Rovira (1959) studied the effect of light and temperature on root exudation. In the wheat rhizosphere, Rouatt and Katznelson (1960) observed that reduction in different bacteria coincided with the reduction of light from 1,000-300 ft.c. However, Peterson (1961) stated that shading of plant<sup>s</sup> had no appreciable effect on vegetatively active fungi colonizing the primary root of wheat and soybean seedlings. Srivastava (1971) studied the effect of continuous light and continuous dark<sup>ness</sup> on the rhizosphere and rhizoplane microflora of wheat and barley. <sup>They</sup> and <sup>he</sup> concluded that amino acids in root extracts were higher in plants exposed to continuous light corresponding with higher microbial population<sup>s</sup> in the rhizosphere<sup>s</sup> of these plants.

*should be scientific writing - content*  
A detailed study of temperature effect has been carried out by Rauatt et al. (1963) with wheat and soybean grown at three ranges of temperature (i.e. 55-60°, 70-75° and 85-90°F). Number<sup>s</sup> of bacteria in the rhizosphere and rhizoplane of wheat increased as the temperature decreased, whereas numbers in the root free soil and on soybean roots increased with increased temperature. Fungal isolation showed a higher incidence of

Mucor, Rhizopus, Rhizoctonia and Gliocladium spp. on soybean at high temperature whereas species of Fusarium and Cylindrocarpon were prevalent at low temperature. However, the most most striking feature was the predominance of non-sporing hyaline type at low temperature. The results suggested that temperature exerted both direct and indirect effect on the rhizosphere population. The later being more important. Namoradize (1975) tabulated effect of different temperatures on the development of Fusarium <sup>spp.</sup> species isolated from peach roots where <sup>cardinal</sup> minimum temperature<sup>s</sup> for mycelial growth of F. oxysporum and F. solani <sup>were</sup> <sup>minimum</sup> ~~was~~ <sup>10°</sup>, optimum 25-30° and maximum 35°C. Sporulation was observed only at 15-30°. Spore germination occurred at 5-37° with optimum temperature of 20-30°C. It is suggested that no general<sup>g</sup>ization can be made with respect to occurrence of different fungi as influenced by temperature. But large <sup>numbers</sup> ~~no~~ of fungi are found at temperatures at which the growth of the plant is favoured (Marks, et al., 1975; Bissett and Parkinson, 1979, and Phillips, 1986).

#### 1.2.8 Seasonal Variation

The primary factor responsible for rhizosphere phenom<sup>the</sup>enon is plant itself. So any condition that significantly effect<sup>s</sup> its morphology and physiology will be reflected in <sup>the</sup> quantitative and qualitative changes in the microflora of root zone. In addition to various effects of stage of the plant

growth and kind of the plant, the percentage of moisture content, reaction and fertility of the soil, environmental factors such as light, temperature and seasonal variations, and plant treatment <sup>a</sup>effect the rhizosphere population to a varying degree by influencing the plant growth. They also exert a direct though considerable less effect on micro-organism themselves. Seasonal effects as reported by various workers (Katznelson, 1946; Millar and Boothroyd, 1962) may be considered to reflect the combined influence of all these environmental conditions. Panwar et al. (1967) observed that Rhizopus, Aspergillus, Penicillium, Chaetomium, Neocosmospora, Alternaria, Helminthosporium, Curvularia and Fusarium were some of the common genera which occurred practically all the year round, while the activity of certain forms like <sup>species of</sup> Choenophora, Cunninghamella, Sordaria, Monocillium, Myrothecium, Cladosporium, Cylindrocephalum, Cephalosporium, Trichothecium sporotrichum and Phoma varied with the plant and the season. Marks et al. (1975) studied quantitative variation in population levels of Phytophthora cinnamoni in Euclyptus <sup>a</sup>forst soil<sup>s</sup> of Eastern Victoria. There was marked seasonal variation in the Population Density Index (PDI), especially on the <sup>??</sup>(hight)hazard site. Minimum values occurred in winter and maximum in summer. Gangawane and Deshpande (1977) observed qualitative and quantitative differences in the rhizosphere mycoflora of groundnut in response <sup>to</sup> of seasonal variations. Davet et al. (1980) studied rhizosphere

mycoflora of bean (<sup>olus</sup>~~Phaseolus~~ vulgaris) and observed that Fusarium solani occurred through<sup>out</sup> the year while Rhizoctonia solani was favoured by warm<sup>temperatures</sup> and Pythium aphanidermatum by <sup>the</sup> rainy season. Similar seasonal variations in the occurrence of fungi in <sup>the</sup> rhizosphere has been reported by Padma and Mukerje, 1972; Subramaniyan and Rao, 1977; Bisset and Parkinson, 1979; Manoharachary and Bilolikar, 1979; Singh and Bhargawa, 1981 and Vardavakis, 1986.

#### 1.2.9 Soil Amendment

Organic amendment of soil has been a village practice for a long time without realising its implication in disease control. Clark and Thom (1939) reported that rhizosphere microbial population of cotton and wheat responded favourably to organic manuring. There <sup>have</sup> ~~has~~ been other reports of increases in microbial population<sup>s</sup> due to organic amendment<sup>s</sup> (Absalyamova, 1963; Bumbieris, 1969; Khanna and Singh, 1975; Ashraf, 1981 and Ansari, 1983). Venkateson (1962) pointed out that, although initially there <sup>was</sup> ~~(has been)~~ a decrease in rhizosphere population of bacteria and fungi in <sup>a</sup> rice field due to organic amendments, <sup>the</sup> (but) after 45 days <sup>R:S</sup> ratio increased. Not only organic amendment but incorporation of fertilizers<sup>also</sup> has resulted in an increase in the microbial counts (Mosolov et al., 1959; Gadzhieva, 1959; Jalaluddin, 1975 and Ansari, 1983). Samtsevich and Borisova (1961) pointed out that, although in pot experiments

soil amendment with mineral and organic fertilizers had little effect on microbial counts in uncropped soil or in wheat rhizosphere, but in field experiments there had been a significant increase in the rhizosphere population which (however) depended on the time of the year. Mishra (1972) recorded a stimulation in the rhizosphere mycoflora of Oryza sativa L. as a result of incorporation of superphosphate. Bagyaraj and Rangaswami (1967), on the other hand, recorded a decrease in population of fungi in the rhizosphere of Eleusina coracana Gaertn due to amendment with fertilizers. At the other extremes, Voroshilova (1956), Katznelson et al. (1959), Lochhead (1959) and Baker and Nash (1965) observed that rhizosphere population was not affected by soil amendments.

This principle of modification of rhizosphere mycoflora has been employed in controlling (the) disease (Davey and Papavizas, 1960 and Singh et al., 1985). Davey and Papavizas (1959 and 1960) pointed out that addition of organic materials (although) resulted in an increase in microbial population but suppressed the root rot of beans caused by Rhizoctonia spp. Chauhan (1963) observed a reduction in the incidence of Fusarium wilt of gram (Cicer arietinum L.) by amending the soil with groundnut, till (Sesamum indicum) or mustard oilcake. Srivastava and Sinha (1971) reported a reduction in wilt of coriander (Coriandrum sativum L.) caused by Fusarium oryso<sup>xy</sup>sporum f.

( )

sp. cCoriandri Narula and Joshi by soil amendment with oilcakes. Soil amendment with chitin, Laminarin, wheat straw and over-dried green clover has been reported to inhibit the microbial population in soil and activity of Verticillium dahliae (Jarden et al., 1972). Population of Fusarium (as a result of organic amendments) was reduced by volatile degradation products of oilseed amendments (Zakaria et al., 1980). Fungistat<sup>c</sup>is' effect against pathogenic fungi has been reported due to soil amendment with mesocarp, root, seed and skin extract of avocado trees (Wehner, 1982) and with N and chitin against Fusarium oxysporum f. sp. ciceri (Satyaprasad and Rao, 1982). There has been <sup>q</sup>reduction in the population of sclerotia of Corticium sasakii (rice pathogen) due to amendment of soil with dhaincha (Sesbania sp.) followed by sunnhemp and mung (Dath, 1982). Similar<sup>1</sup> reduction<sup>5</sup> in population<sup>5</sup> of soil borne pathogens have been reported by Lumsden (1983) and Sinha and Prasad (1986). (There have been report where<sup>1</sup> incidence<sup>1</sup> of diseases <sup>also have</sup> had<sup>1</sup> been found to increase due to soil amendments. Son et al. (1985) reported that incidence of Ginseng root rot caused by Fusarium solani increased due to soil amendment with (amendments of) crushed wheat, sweet potato stalk, chinese cabbage, ginseng leaves and soybean pods, but soil amendment with garlic roots, walsh onion, cabbage leaf and stalk, green onion stalk, wheat straw and barley straw decreased the incidence. Similarly Moubasher and Abdel Hafez



(1986) pointed out that there had been stimulation of rhizosphere fungi of cotton and root pathogens (i.e. Rizoctonia solani, Fusarium oxysporum and F. moniliformae (Gibberella fujikuroi)) by dung manure, clover straw and wheat straw.

O'Rourke and Miller (1966) pointed out that soil amendment with fertilizers decreased the incidence of alfalfa root rot and recovery of Fusarium spp. from roots and rhizosphere. Reduction in the number of <sup>propagules of</sup> Fusarium oxysporum f. sp. niveum (wilt pathogen of watermelon) due to <sup>soil</sup> amendment with N fertilizers, including urea and  $\text{KNO}_3$ , and increases <sup>in population</sup> with ammonium sulphate have been reported by Huang and Sun (1982). Addition of urea in soil increased the amount of ammonia which resulted in the decrease in the population of Pythium ultimum, Thielaviopsis basicola and Macrophomina phaseolina (Lockwood, 1985). The population of Acremonium implicatum and Fusarium sacchari was <sup>? reduced?</sup> recorded by soil amendment with inorganic substances (Narendra et al., 1985). This was attributed to be due to increase in soil populations of antagonistic and fungistatic actinomycetes, <sup>other</sup> bacteria and fungi.

Inhibition of soil microorganisms and plant pathogens due to application of pesticides, herbicides, fungicides and nematicides has also been reported (Bollen et al., 1954; Bollen, 1961; Lebed, 1964; Bazhina, 1967; Alconero and

Ragedorn, 1968; Valaskova, 1968 and 1969; Smiley et al., 1970; Agnihotri, 1971; Chawdhery et al., 1972, Tu, 1972 and 1973; Singh and Prasad, 1973; Maric et al., 1974; Midhu and Nandwana, 1974; Phipps and Stipes, 1975; Jardon, 1975; Attabhanyo and Holcomb, 1976; Bertoldi et al., 1977 and 1978; El-Khadem et al., 1977; Oku, 1979; Sinha et al., 1980; Jain and Sehgal, 1980; Bartaria et al., 1982; Reis, 1982; Bollen et al., 1983; Malajczuk et al., 1983; Doneche et al., 1983; Torststensson and Wessen, 1984; Deb and Dutta, 1984; Ingham, 1985; Thind and Jhooty, 1985; Youseff et al., 1985; Yarden, 1986). This reduction has been reported in part due to direct influence on <sup>the</sup> pathogen <sup>and</sup> also in parts due to disturbance of soil microflora.

#### 1.2.10 Foliar Spray

Besides soil amendments, foliar spray of plants with chemicals, which could be pesticide<sup>s</sup> or fertilizer<sup>s</sup>, also influence the rhizosphere microflora, both qualitatively and quantitatively (Rovira, 1959, Rauatt and Katznelson, 1960; Balasubramanian and Rangaswami, 1973 and 1975). Ramachandra Reddy (1959 and 1968) observed a quantitative and qualitative change in the population of bacteria, actinomycetes and fungi in the rhizosphere of rice by spraying with urea. There <sup>was</sup> ~~has been~~ a greater influence in the number of Penicillia <sup>spp.</sup> ~~at low~~

Can not use this term.  
Only one species of  
"Penicillium" recognized.  
Can not have plural penicillia  
rather use penicillium spp.  
(for plural species). Confirmed

concentration<sup>s of urea</sup> but at higher concentration<sup>s</sup> of urea Aspergillus <sup>s.s. 11</sup> were stimulated. Horst and Herr (1962) reported an increase in the number of actinomycetes antagonistic to Fusarium roseum f. cerealis in the rhizosphere due to spray<sup>s</sup> of plants with urea. Similar increase<sup>s</sup> in bacteria and ant<sup>e</sup>gonistic actinomycetes due to spray<sup>s</sup> with urea <sup>have</sup> ~~has~~ been observed by Vransy (1972). As a result of spray<sup>s</sup> with urea, there has been an increase in amino acid content, particularly glutamic acid and  $\alpha$ -amino butyric acid in the root exudates of plants (Agnihotri, 1964). Spray<sup>s</sup> of plants with ammonium sulphate increased the population of fungi and actinomycetes (Bagyaraj and Rangaswami, 1982). On the other hand, a decrease in the number of fungi in the rhizosphere<sup>s</sup> of wheat and maize as a result of spray<sup>s</sup> has also been observed (Vransy, 1963 and Annapurna and Rao, 1983).

Changes in the population of microorganisms in the rhizosphere as a result of sprays with other chemicals such as antibiotics, nutrients and growth regulators have also been recorded (Venkatasana, 1960, Rangaswami and Vasanthrajan, 1961; Vransy et al., 1962; Kundaswamy and Rangaswami, 1967; Sullia, 1968; Gujrati, 1969; Singh, 1970; Gupta, 1971 and Chandra et al., 1982). Dwivedi and Singh (1971) noted that spray<sup>s</sup> with gibberelllic acid in low concentration (i.e. 50 ppm) increased the fungal population but spray<sup>s</sup> with higher concentrations

(100 and 200 ppm) inhibited the buildup. However, maleic hydrozide was found to be inhibitory at all the concentrations tested.

*Some*  
✓ ~~fungicides~~ fungicides sprayed for controlling diseases are transported downwards into roots. Thus the root exudates are also influenced, which in turn influence the rhizosphere microflora. There ~~has been~~<sup>was</sup> a decrease in the rhizosphere population of bacteria as a result of spray with Bordeaux mixture, malachite green and Dithane Z-78 (Halleck and Cochrane, 1950) and with other fungicides (Swaminathan and Sullia, 1969 and Wainwright and Sowden, 1977).

Balasubramanian and Rangaswami (1973) pointed out that application of Diathane Z-78 (200 ppm) adversely influenced the exudation of amino acids from the roots of Sorghum vulgare and Crotolaria juncea L., which probably caused reduction in rhizosphere population of microorganisms. Similar results were obtained by Sullia (1969), Srivastava and Mishra (1971), Ranga Rao et al. (1972), Wainwright and Sowden (1977), Srivastava and Dayal (1981) and Gunasekar and Rao (1982).

#### 1.2.11 Pathogenesis and Resistance

The intense microbial activity in the rhizosphere has direct or indirect effect on plant pathogen<sup>s</sup> and development of diseases (Skilyagina, 1967; Voronkevich et al., 1972;

Mall, 1973; Wnekowski, 1975; Kutrzeba, 1984; Gagne et al., 1985 and Sakthivel et al., 1986) and vice-versa (Parlik, 1950; Agnihothrudu, 1959; Mathur and Chauhan, 1972; Stefurak, 1976; Ellingboe, 1983 and Yuen and Schroth, 1986).

By and large, the rhizosphere microflora constitute the non-pathogenic organisms. This often leads to more destructiveness <sup>caused by</sup> of soil borne pathogens in sterile than in normal, <sup>non-</sup>unsterilized soil. Even susceptibility and resistance to soil borne pathogens have been related to the activity of soil/rhizosphere microflora. Reynolds (1926) correlated the resistance of flax varieties to Fusarium lini (Bolley) to the availability of hydrocyanic acid. In the plants, a greater rhizosphere activity in the susceptible varieties has been reported in (many) crops <sup>such as</sup> (eg.) flax against Fusarium oxysporum and tobacco against Thielaviopsis basicola Zopt. (Timonin, 1940), banana (Harper, 1950), maize (Horny and Ullstrup, 1967), cotton (Avezdzhanova, 1974), wheat (Srivastava and Mishra, 1972) and cauliflower (Khan et al., 1973). Species of Alternaria, Cephalosporium, Fusarium, Helminthosporium and Verticillium were more abundant in susceptible varieties of flax as compared to resistant varieties (Timonin, 1941). Winter and Rumker (1949) <sup>concluded</sup> on the basis of detailed studies ~~(concluded)~~ that resistance of maize and wheat to Aschochyta pinodella Johns and Fusarium culmorum (Smith) Saccardo respectively was mostly conditioned by rhizosphere microflora.

Parlik (1950) observed that Aspergillus glaucus Link and related species predominated in the sterile soil, but Fusarium lini Bolley and Penicillium spp. <sup>predominant</sup> in the rhizosphere soil of flax seedlings infected with F. lini. (There has been) <sup>high</sup> rate of multiplication of species of Penicillium in the rhizosphere of infected plants, but the multiplication of <sup>occur</sup> saprophytic bacteria and fungi was markedly reduced. Species of Aspergillus, Phoma, Pythium, Rhizoctonia <sup>U</sup> were found predominating in the rhizosphere of diseased plants of Pinus <sup>sp.</sup> while <sup>sp.</sup> of Alternaria, Cephalosporium, Metarrhizium, Spicaria and Tilachlidium spp. <sup>predominant</sup> in healthy plant <sup>S</sup> (Timonin, 1966). Buxton (1957) reported that germination of conidia of a race of Fusarium oxysporum f. <sup>sp.</sup> pisi Linford was poor in the extract of the rhizosphere of resistant cultivar <sup>S</sup> but was stimulated in that of susceptible cultivar <sup>S</sup>. This led him to conclude that rhizosphere microflora <sup>are</sup> (is) affected by the excretions from the roots of susceptible varieties. Similar results have been obtained with varieties of Gossypium arborium race indicum lini (Sulochna, 1958).

Mukhopadhyay and Nandi (1974) reported, differences between healthy jute plants and plants infected with Macrophomina phaseoli (Maubl.). Agnihothrudu (1959), while studying the rhizosphere microflora of tea in relation to charcoal stump rot caused by Ustilina zonata (Lev) Sacc., pointed out that

there were differences in density of the fungi and <sup>b</sup>~~acteria~~ in rhizosphere flora of apparently healthy and infected plants. Khare (1968) recorded that the rhizosphere of diseased strawberry plants harboured more fungal population in comparison to their healthy counterparts. Powell (1969) observed that Trichoderma sp. exhibited higher frequency in healthy plants and not in infected ones.

Vishwanath et al. (1969) pointed out that the microbial population<sup>s</sup> of soil and rhizosphere ~~was~~ <sup>were</sup> altered both qualitatively and quantitatively around diseased coffee plants. Mathur and Chauhan (1972), while studying the rhizosphere of gram infected with Fusarium oxysporum f. sp. ciceri Natuo sato., Rhizoctonia solani Kuhn and Sclerotium <sup>roissii</sup> roffsii Sacc., recorded higher population<sup>s</sup> of microorganisms in the rhizosphere of diseased plants. Similar results have been obtained by Srivastava and Mishra (1972) in wheat infected with Puccinia graminis var. tritici (Pers.) Eriks and Henu: by Khan et al. (1973) in cauliflower infected with Rhizoctonia solani (Kuhn) and Fusarium spp.; by Babushkina (1973) in cotton infected with Verticillium dahliae Kleb; by Rai and Upadhyay (1980) in pigeonpea infected with Fusarium udum; by Ashour et al. (1980); in onion infected with (Fusarium oxysporum f. sp. cepae; by Ashraf (1981) in Triticales hexaploid and Pennisetum typhoides infected with Puccinia graminis and Sclerospora sp. respectively, and by

Wadhvani and Mehrotra (1982) in smut infected plants of Cynedon dactylon. Vesely (1985) concluded that the numbers <sup>were higher</sup> of pathogenic and saprophytic fungi ~~(was)~~ <sup>more</sup> on the root surface of sugarbeet plants affected by damping-off. On the other hand, Chatopadhyay and Mukhopadhyay (1967) recorded higher rhizosphere population in healthy rice plants in comparison to their infected counterparts. Similar results were obtained by Stefurak (1976) in spruce infected with Fomitopsis annosa Karst (Heterobasidium annosum), by Kannaiyan and Prasad (1981) in rice infected with Rhizoctonia solani and by Kishan et al. (1982) in sugarcane infected with Glomerella tucumensis. In multiple pathogenic conditions, such as root knot and Fusarium wilt, there has been greater microbial activity in the rhizosphere of plants. Bergeson et al. (1970) in disease complex studies concluded that Fusarium oxysporum f. sp. lycopersici was more in the rhizosphere of tomato infected with Meloidogyne javania than in rhizosphere of non-infected plants. Hirano (1983) reported that in multiple pathogenic conditions involving root-knot nematode and Fusarium sp. in tomato there <sup>was</sup> ~~has been~~ an increase in the population of gram <sup>positive</sup> bacteria but reduction in Fusarium oxysporum propagulas. <sup>gram negative</sup>

### 1.2.12 Interactions Between Soil Microorganisms

Different rhizosphere fungi develop some kind of relationship with each other. (The most important one is the antagonistic

Omit sentence unless you sit proof of importance. Beneficial relationships may be more important.



relationship. Waksman (1944) established the existence of mycelia in soil and hence suggested active involvement of fungi in the soil ecosystem. Soil fungi such as Chaetomium spp. had an antagonistic effect on <sup>species</sup> Aspergillus, Curvularia, Fusarium and Penicillium (Mukerji, 1968). Trichoderma lignorum (T. viridae), T. koningi and Pseudomonas fluorescens, Bacterium pyocyaneum (P. aeruginosa), Bacillus subtilis, B. megaterium and B. mesentericus (Anon, 1968). Babushkina (1974) pointed out that there <sup>have</sup> ~~has~~ been more antagonistic fungi in the rhizosphere of healthy plants than that of Verticillium-infected plants. Similar results were obtained by Mall (1973). Penicillium carpeteales, Chaetomium and Trichoderma have been found antagonistic to various pathogens such as Fusarium moniliforme (Gibberella fujikaroi) (Toledo and Cardoso, 1975) Trichoderma spp. to Phytophthora (nicotiana var.) parasitica, Rhizoctonia solani, Fusarium oxysporum and Verticillium dahliae (Raicu and Stan, 1975), Trichoderma harzianum to Sclerotium rolfsii (Arora and Dwivedi, 1979), Corticium rolfsii, Fusarium oxysporum f. lycopersici (Endo et al., 1976). Pineda and Diaz-Palanco (1981) observed reduction in the incidence of Sclerotium rolfsii on bean by Penicillium notatum. Dwivedi and Mishra (1982) observed that Cladosporium cladosporioides was a hyperparasite on Rhizopus oryzae. Vargas and Ramirez (1983) reported that Bacillus megaterium from the rhizosphere of cotton seedlings strongly inhibited Rhizoctonia solani,

the causal agent of damping-off. Lumsden (1983) established effective biological control of Sclerotium spp., Sclerotinia spp., Rhizoctonia solani and Pythium spp. by Trichoderma spp. and Gliocladium spp.; Sclerotinia spp. and Sclerotium cepivorum by Sporidemia sclerotivorum and Coniothyrium minitans, Pythium spp. and Rhizoctonia solani by Tisaria arvalis and Fusarium spp. and Pythium spp. by Chaetomium globosum. Gokulopalan and Nair (1984) pointed out that Aspergillus niger and Trichoderma viridae were highly antagonistic to Rhizoctonia solani. Hus and Lockwood (1984) observed that Hyphochytrium catenoides could be used for biological control of Phytophthora root rot. Aspergillus niger has also been reported to be antagonistic to the collar rot pathogen, Rhizoctonia solani (Venkatasubhaiah and Safeulla, 1984). Trichoderma spp., Cynothyrium minitans and species of Fusarium and Penicillium isolated from soil have also been shown to have antagonistic effect<sup>s</sup> on Sclerotinia sclerotiorum (Zozzerini and Tosi, 1985). Acrophyalophora levis, Myrothecium roridum and Paecilomyces inflatus exhibited strong antagonism<sup>s</sup> towards B. theobromae; Aspergillus niger and Memmoniella echinata towards F. solani; and Aspergillus niveus and Mimmoniella echinata towards Macrophomina phaseolina (Agarwal, 1986). Trichoderma harzianum, Penicillium purpurogenum and P. funiculosum were found to be the most antagonistic to Drechslera sorokiniana (Cochliobolus sativus) (Ermeкова and

This is not referring to the work cited for Lumsden. You must refer the work cited here to the original paper. It is also interesting to know if lower said in the case of green house and in some cases not in the soil and not sufficient bio logical control!! et al.

but works in greenhouse!

Abirova, 1986). Thus, it is clear that there are a large number of fungi which are antagonistic to pathogenic fungi and the most important one <sup>may be</sup> is species of Trichoderma.

Legumes, <sup>which are</sup> ~~being next to cereals is~~ <sup>are</sup> grown the world over as source<sup>s</sup> of proteins (Allen and Allen, 1958) and form an important constituent of vegetarian diets. Besides protein, they are also source of carbohydrate, fats, vitamins and minerals.

Chickpea (Cicer arietinum L.) is the third <sup>most</sup> important pulse crop in the world from extending latitude from 15 to 40 N of equator. In India, the area under cultivation of chickpea is 9 million hectares (Chandra, 1983) and <sup>provides</sup> ~~amount~~ 90% of the world production (Verma, 1984). The yield of the crop in India is far below the average yield of chickpea in developed countries, which is partly due to poor tillage practices and partly due to damages done by diseases and pests (Pandey, 1982; Tripathi and Sharma, 1983, Singh and Bhargawa, 1985 and Grewal et al., 1985). Among the various disease<sup>s</sup> of chickpea, wilt<sup>s</sup> occupy an important position in lowering the yield of crop.

The wilt caused by Fusarium oxysporum Schlecht. emend. Snyd. and Hans. f.sp. ciceri (Padwick) Snyd. and Hans. was first described by Padwick in 1940 from India. Annual losses to crop due to the disease <sup>have</sup> ~~has~~ been estimated to be 10 percent (Haware and Nene, 1982). The causal organism is both seed

borne and soil borne (Haware et al.; 1978; Kotasthan et al., 1979; Sharma and Gupta, 1983; and Conci et al., 1985).

Soil borne diseases have been controlled by managing the rhizosphere mycoflora favouring the antagonists of the causal organism in the soil (Winter and Rumker, 1949; Babushkina, 1974; Vorgas and Ramirez, 1983 and Zozzerini and Tosi, 1985). This has been successfully attempted by comparing the rhizosphere mycoflora of infected and uninfected crop followed by management of antagonists by foliar sprays and organic amendments (Davey and Papavizas, 1960 and Horst and Herr, 1962; Vransy, 1972 and 1973; and Bagyaraj and Rangaswami, 1982 and Narendra et al., 1985).

From the forgoing review of literature it is clear that very little work has been done on the rhizosphere and rhizoplane mycoflora of gram in relation to wilt disease caused by Fusarium oxysporum f. ciceri. Moreover, little information exist <sup>as to whether or not</sup> ~~if~~ this can be controlled by modifying the rhizosphere mycoflora. Hence in the present investigations, an attempt has been made to study the following:

1. Rhizosphere and rhizoplane mycoflora of different cultivars of chickpea inoculated with F. oxysporum f. ciceri and compared with <sup>those of</sup> uninoculated plants.
2. Rhizosphere and rhizoplane mycoflora of chickpea inoculated with F. oxysporum f. ciceri at varying age<sup>s</sup> and compared with

those of  
uninoculated plants.

3. Effect of spray<sup>ing plants</sup> with indole acetic acid, indole butyric acid, thio-indole butyric acid, gibberellic acid, maleic hydrazide, urea, potash, bavistin, vitavax, brassicol, benlate, captan, wettable sulphur and streptomycin on rhizosphere and rhizoplane mycoflora of cultivar JG-62 and JG-74 inoculated with F. oxysporum f. ciceri (and compared with uninoculated plants).
4. Effect of soil amendments with urea, superphosphate, potash, castor cake, mustard cake, neem cake, mahua cake, bavistin, vitavax, brassicol, benlate, captan, wettable sulphur on rhizosphere and rhizoplane mycoflora of chickpea cultivars JG-62 and JG-74 inoculated with F. oxysporum f. ciceri (and compared with uninoculated plants).

Refer to JG-62 and JG-74  
which are not inoculated  
with F. oxysporum f. ciceri  
in the above experiments.

## CHAPTER - II

### MATERIALS AND METHODS

*These need to be summarised, especially in rhizosphere fungi. While treatments, since as they have shown these fungi affect rhizosphere, not.*

#### 2.1 PREPARATION OF FIELD

Plants were grown in beds each with an area of 10 sq. meters and were given routine agronomic practices. The beds were arranged in randomised fashion and each treatment was replicated thrice.

#### 2.2 ISOLATION OF FUNGI FROM RHIZOSPHERE, RHIZOPLANE AND NON-RHIZOSPHERE

For isolating fungi from rhizosphere, five plants were removed carefully from the field and brought to the laboratory in sterilized containers under aseptic conditions. Blocks of soil adhering the roots were cut and crushed gently with minimum damage to the roots. The roots were shaken gently to remove the superfluous soil and were transferred <sup>into a</sup> ~~in the~~ beaker containing 100 ml of sterilized distilled water and shaken. To 10 ml of this suspension 90 ml of sterilized distilled water was added in a separate container. Further, 1 ml from this solution was transferred to in another flask containing 99 ml of sterilized, distilled water. Each time the flask was shaken.

*40 ml of this suspension was added in a separate container.*

In this way <sup>a</sup> suspension of 1:10,000 was obtained. With the help of a sterilized pipette, 1 ml of <sup>the</sup> 1:10,000 dilution was placed <sup>each 2X</sup> in sterilized petridishes along with 10 ml of melted cooled peptone dextrose rose bengal agar<sup>+</sup>.

<sup>that</sup> The fungi developed after one week were examined and identified. The frequency was calculated by the formula:

$$\frac{\text{Number of plates containing a particular fungus}}{\text{Total plates poured}}$$

For determining the weight of rhizosphere soil, roots were removed from the original dilution flask and washed. The washed water was collected in the original flask containing stock solution. The water was evaporated on a waterbath and the soil residue was dried to constant weight in an oven at 105°C. The flask containing dry soil was weighed and the dilution factor calculated, allowance being made for the amount of soil removed in preparing the dilutions. Plate counts were made and the number of fungi per gram of oven dry "rhizosphere soil" was calculated.

---

<sup>+</sup>PEPTONE DEXTROSE ROSE BENGAL AGAR MEDIUM

Agar	20.0 g
KH <sub>2</sub> PO <sub>4</sub>	1.0 g
Peptone	5.0 g
Dextrose	10.0 g
Distilled H <sub>2</sub> O	1000.0 ml
Rose bengal	1:30,000
Streptomycin	30 µg/ml (Martin, 1950; Johnson, 1957)

For isolating rhizoplane mycoflora the "Serial root washing technique" of Harley and Waid (1955) was employed. Roots were washed several times in sterilized distilled water, and cut into small pieces of 3 mm length, and transferred into sterilized petriplates, with five root pieces in each petriplate containing 10 ml sterilized, melted, cooled peptone dextrose rose bengal agar medium. The fungi that developed were examined and identified. The frequency of the fungi was calculated by the method given earlier.

For studying the non-rhizosphere soil mycoflora, soil was brought from uncultivated portions of the field in a randomised manner from a depth of 10 cm. Under aseptic conditions, the samples were mixed thoroughly in order to get a composite sample. Twenty five g of soil was taken from this composite sample and transformed in 250 ml of sterilized distilled water. Flask was shaken vigorously and the resultant suspension was used as a stock solution. From this stock solution a dilution of 1:10,000 was achieved by making transfers in the manner given earlier. One ml of 1:10,000 dilution was transferred into petriplates along with 10 ml of melted cooled peptone dextrose rose bengal agar medium. Petriplates were rotated gently in order to obtain an equal distribution of solution and medium. The fungi which developed were identified and counted. The frequency of the fungi was calculated by the





The latter, <sup>5</sup>fungus was transferred into 4 holes made around the roots of 15 day old seedlings.

## 2.5 BIOCHEMICAL ANALYSIS OF ROOT

### 2.5.1 Preparation of Root Powder

Roots of both <sup>non-</sup>~~un~~inoculated and inoculated plants (25 plants each) were collected from the field and were washed thoroughly in running water and then in sterilized distilled water. These roots were dried in an incubator (~~running~~) at 60°C for 48 hours. and powdered in an electric grinder. The powder was sieved through<sup>a</sup> 60-mesh sieve.

### 2.5.2 Quantitative Estimation of Total Free Amino acids

Amino acids were estimated ~~(by)~~ using the method of Moore and Stein (1954) <sup>with</sup> ~~using~~ modified ninhydrin reagent. Ninhydrin reagent was prepared by dissolving 20 g of ninhydrin and 3 g of hidrindentine<sup>+</sup> in 750 ml of methyl cellosolve (2-methoxy-ethanol). The mixture was gently stirred. To this <sup>mixture</sup> 250 ml of

HYDRINDENTIN: 8 g of ninhydrin was dissolved in 200 ml of distilled water at 90°C. In another solution 8 g of ascorbic acid was dissolved in 40 ml of distilled water at 40°C. The two solutions were immediately mixed and later cooled under tap water. The crystals of hydrindentin thus obtained after filtration were washed and dried in a vacuum dessicator in dark and stored in dark glass bottle.

sodium acetate buffer<sup>+</sup> of pH 5.5 was added. The resulting reddish solution was immediately transferred to a dark glass bottle.

For the preparation of<sup>a</sup> standard curve, 1 mg of leucine was dissolved in 100 ml of double distilled water. One ml of this solution was diluted to 10 ml by adding<sup>the</sup> required amount of water. Different (~~amounts of~~) aliquotes<sup>of</sup>, 0.1, 0.2, 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, 0.9 and 1.0 ml were taken and the final volume in each case was made to 1 ml by further addition of distilled water. To 1 ml of the above solution 1 ml of<sup>in 1 ml of</sup> ninhydrin reagent was added and the solution was passed<sup>in 1 ml of</sup> for 15 minutes over a water bath. The violet coloured solution was diluted to 10 ml by adding more distilled water. Optical densities were measured in<sup>a</sup> Bousch and Lomb Spectronic 20-Calorimeter at 570 nm against the reagent blank. A graph between<sup>of</sup> the optical densities and different concentrations was plotted, which was a straight line.

---

<sup>+</sup>SODIUM ACETATE BUFFER: The 4 N sodium acetate buffer of pH 5.5 was prepared by dissolving 272 g of sodium acetate (reagent grade) in 700 ml of distilled water over<sup>a</sup> waterbath. After cooling at room temperature 50 ml of glacial acetic acid was added and then<sup>a</sup> final volume was made up to 500 ml. The pH of the buffer was later checked by pH meter.

For the estimation of total free amino acids, 100 mg <sup>of</sup> dried sample of powdered material <sup>were</sup> ~~was~~ extracted with 5 ml <sup>of</sup> 80 percent boiling ethanol and centrifuged at 3000 rpm for 15 minutes. <sup>The</sup> ~~The~~ Extract thus obtained was diluted to 10 ml by adding <sup>the</sup> ~~the~~ required amount of 80% ethanol. To 1 ml of this solution, 1 ml ninhydrin reagent and 8 ml distilled water were added and the optical densities were determined.

### 2.5.3 Qualitative Estimation of Amino acids

Qualitative analysis of amino acids was done by following the method of Block (1950) using <sup>a</sup> ~~a~~ descending chromatography with Watman filter paper no.1 and n-butanol:glacial acetic acid:water (12:3:5) as solvent. Alcoholic extracts of samples along with 22 known amino acids were spotted on one end of the sheet in a line with the help of <sup>a</sup> ~~a~~ micro pipette and suspended. After 10 hours, the sheets were dried in the air and sprayed with 0.2% (v/v) of ninhydrin in water saturated n-butanol. The sheets were dried in air followed by drying in <sup>an</sup> ~~an~~ oven at 100°C for 2 minutes. The R/S value of various spots developing were measured and were compared with R<sub>f</sub> values of known amino acids.

R<sub>f</sub> value = a/b, where a = distance travelled by the known sample in cm and b = distance travelled by the solvent in cm.

#### 2.5.4 Estimation of Phenols

Phenols were estimated following the method of Biehn et al. (1968) using Folin-Ciocalteu reagent (Bray and Thrope, 1954). Folin-Ciocalteu reagent was prepared by dissolving 100 g of sodium tungstate and 25 g of sodium molybdate in 700 ml of distilled water. To the above solution 50 ml of 85% phosphoric acid and 100 ml of concentrated hydrochloric acid were added. This was then refluxed for 10 hours. Later, 150 ml of Lithium sulphate, 60 ml of double distilled water and few drops of liquid bromine were added. The mixture was then boiled over a free flame for about 15 minutes to remove excess bromine and was cooled at room temperature. The solution was filtered and <sup>adjusted</sup> ~~made~~ to 1000 ml by adding more distilled water. Its normality was adjusted to 1-N before use.

A standard curve of different concentrations of phenol was prepared by dissolving 10 mg of paracresol in 100 ml of 80% butanol. One ml of paracresol solution was diluted again by adding 10 ml of 80% ethanol. From this solution, 0.1, 0.2, 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, 0.9 and 1.0 ml were transferred to test tubes and the volume in each case was made up to 1 ml by adding <sup>the</sup> required amount of 80% ethanol. Two drops of concentrated hydrochloric acid were added and then the solution was heated over flame, avoiding overheating. Thereafter, 1 ml of 1-N Folin-Ciocalteu reagent and 20 ml of 4% of sodium carbonate were

added. After 20 minutes the optical density was measured by using <sup>a</sup>Bausch and Lomb Spectronic-20-Calorimeter at 660 nm against a reagent blank. A straight line curve, obtained by plotting different concentration<sup>s</sup> of paracresol and optical densities, served as standard curve.

For estimating total phenols in the sample, 100 mg dried, powdered sample <sup>were</sup> was heated with 5 ml of 80% ethanol for 15 minutes over <sup>a</sup>waterbath; <sup>the suspension was</sup> and then centrifuged for 5 minutes at 3000 rpm. The alcoholic extract was taken out and made up to 10 ml by addition <sup>of the</sup> required amount of 80% ethanol. To 1 ml of this solution, 1 ml Folin-Ciocalteu reagent, 2 drops of concentrated hydrochloric acid and 20 ml of 4% sodium carbonate were added. Optical density was measured and the concentration of phenols was calculated by comparing with the standard curve.

#### 2.5.5 Estimation of O-dihydroxy phenols

Estimation of O-dihydroxy phenols was made following the method of Johnson and Schal (1954) by using Arnows reagent.

The following three stock solutions were prepared:

1. 0.5-N-hydrochloric acid;
2. Nitrate molybdate reagent: 10 g of sodium molybdate <sup>were</sup> was dissolved in 100 ml of distilled water; In this solution 10 g of sodium nitrate <sup>were</sup> was dissolved;
3. 1-N <sup>s</sup>sodium hydroxide.

For obtaining<sup>a</sup> standard curve, 10 mg of catechol ~~was~~<sup>were</sup> dissolved in 100 ml of 80% ethanol. One ml of this solution was again diluted by adding 10 ml of 80% ethanol. (Different) amounts of aliquots<sup>b</sup> 0.1, 0.2, 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, 0.9 and 1.0 ml were taken and the final volume in each case was made to 1 ml by adding<sup>adjuster</sup> required amount<sup>the</sup> of 80% ethanol. To each (of the) 1 ml of the above solution, 1 ml of 0.5 N HCl and 1 ml of molybdate reagent<sup>was</sup> was added. (As a result of which yellow colour developed.) To this<sup>the</sup> yellow coloured solution<sup>that contained</sup>, 1 ml of 1N NaOH was added and the volume was made up to 5 ml by adding<sup>the</sup> required amount of distilled water, which resulted in the change of colour to red. This red coloured solution was read in<sup>a</sup> Bausch and Lomb Spectronic 20-Calorimeter at 530 nm against<sup>a</sup> reagent blank and the optical density was noted. <sup>The</sup> Standard curve was obtained with the help of catechol.

For estimation of O-dihydroxy phenols, 100 mg<sup>b</sup> dried sample of powdered material<sup>was</sup> was extracted with 5 ml 80% boiling ethanol, (and) centrifuged at 3000 rpm for 15 minutes, and the filtrate was collected. The final volume of alcoholic filtrate was adjusted to 10 ml by adding<sup>the</sup> required amount of 80% ethanol. One ml of each of the<sup>a</sup> sample obtained above was<sup>added a</sup> taken to 5 ml calibrated test tubes to which a reagent (containing 1 ml of 0.5 NHCl + 1 ml molybdate nitrate reagent + 1 ml of 1N NaOH + water to the final volume<sup>5</sup> 5 ml) was added and the optical density was noted. The

concentration of total O-dihydroxyphenols was calculated by comparing <sup>sum</sup> ~~the~~ with <sup>the</sup> standard curve.

#### 2.5.6 Estimation of Total Carbohydrates

Total carbohydrates were extracted and estimated by the method of Yih and Clark (1965) and Dubois et al. (1956), respectively.

For making <sup>the</sup> standard curve, 10 mg of glucose <sup>analar</sup> was dissolved in 100 ml of distilled water, and 1 ml of this solution was again diluted to 10 ml by adding <sup>the</sup> required amount of distilled water. Different amounts viz., 0.1, 0.2, 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, 0.9 and 1.0 ml, of this solution were transferred to test tubes and <sup>the</sup> final volume of each was made upto 1 ml by adding <sup>the</sup> required amount of distilled water. To these solutions 1 ml of 5% ethanol and 5 ml of concentrated  $H_2SO_4$  (analar) were added. After 10 minutes, the optical density was measured in a Bousch and Lomb Spectronic 20-Calorimeter at 490 nm as against a reagent blank. A graph was plotted between optical density and different concentrations of glucose which served as standard curve.

For estimation of total carbohydrates, 100 mg <sup>of</sup> dried powdered sample <sup>was</sup> boiled in 5 ml concentrated 1N  $H_2SO_4$  for 30 minutes over <sup>a</sup> waterbath; <sup>the suspension was then</sup> ~~and later was~~ centrifuged at 1000 rpm for 10 minutes. The solution was collected in <sup>a</sup> flask and the



residue was washed twice with distilled water. This aqueous filtrate was added <sup>to</sup> ~~in~~ the filtrate kept in flask and volume was made upto 100 ml by adding distilled water. To 1 ml of this solution, 1 ml of 5% ethanol and 5 ml of concentrated  $H_2SO_4$  was added. Optical density was measured and values of total carbohydrates calculated from the standard curve.

## 2.6 RHIZOSPHERE AND RHIZOPLANE MYCOFLORA OF DIFFERENT CULTIVARS OF CHICKPEA PLANTS UNINOCULATED AND INOCULATED WITH FUSARIUM OXYSPORUM F.SP. CICERI

*Same details of infection*

Seeds of different cultivars of chickpea (viz. JG-62, H-208, BG-309, JG-315, BG-212 and JG-74) were sown in soil infested with F. oxysporum f.sp. ciceri. Studies for rhizosphere and rhizoplane mycoflora were made when the plants had attained the age of 45 days. Estimation of total free amino acids, phenols, O-dihydroxyphenols and sugars and qualitative detection of amino acids were also made from the roots of inoculated plants of different cultivars. Similar studies were made from plants grown in uninoculated beds to serve as control.<sup>S</sup>

*Clarify? Same (control) conditions if any infested soil?*

## 2.7 EFFECT OF AGE OF PLANTS ON RHIZOSPHERE AND RHIZOPLANE MYCOFLORA OF INOCULATED CHICKPEA PLANTS WITH FUSARIUM OXYSPORUM F.SP. CICERI

*How many*

The seeds of cultivar JG-62 were sown after surface sterilization with 1:1000 mercuric chloride followed by

repeated washings with sterilized distilled water. Beds were inoculated with the pathogen F. oxysporum f.sp. ciceri.

Uninoculated beds served as control. Rhizosphere, rhizoplane and non-rhizosphere soil mycoflora studies were made at 15-days intervals from the emergence of seedling to senescence. Quantitative estimation were made for total free amino acids, phenols, O-dihydroxyphenols and sugars from root extract of inoculated and uninoculated plants.

## 2.8 RHIZOSPHERE AND RHIZOPLANE MYCOFLORA OF UNINOCULATED AND INOCULATED PLANTS (WITH FUSARIUM OXYSPORUM F.SP. CICERI) OF CHICKPEA CULTIVARS JG-62 AND JG-74 IN RELATION TO

### FOLIAR SPRAYS

Fifteen-day-old

15 days old plants of JG-62 and JG-74 inoculated with F. oxysporum f.sp. ciceri were sprayed with 100 ppm solutions of indole acetic acid, indole butyric acid, thio-indole butyric acid, maleic hydrazide, streptomycin, urea and muriate of potash; 1500 ppm of bavistin, vitavax, brassicol, benlate, captan and wettable sulphur, (separately). Similar sprays were done on uninoculated plants. In order to be sure that sprays are done on foliage, the surface of the soil was covered with plastic sheets. Plants sprayed with distilled water served as control. Subsequent sprays were done at 15-day intervals from the date of 1st spraying. After 14 days of each spray, the rhizosphere, rhizoplane and non-rhizosphere mycoflora were studied.

Do not start  
as there with  
any one  
with me.

How inoc-  
ulated-  
How many?

2.9 RHIZOSPHERE AND RHIZOPLANE MYCOFLORA OF UNINOCULATED AND INOCULATED PLANTS (WITH FUSARIUM OXYSPORUM F.SP. CICERI) OF CHICKPEA CULTIVARS JG-62 AND JG-74 IN RELATION TO SOIL AMENDMENTS

The soil was amended with urea (11 lb per acre), super-phosphate (20 lb per acre), <sup>Rh-S</sup> muriate of potash (70 lb per acre), neem cake, mustard cake, castor cake (100 lb per acre each); mahua cake (50 lb per acre), vitavax (30 kg per hectare), bavistin, brassicol, bentate, captan and wettable sulphure (6 kg per hectare each). After 15 days, ~~of waiting~~ seeds of JG-26 and JG-74 were sown. Half of the beds were inoculated with F. oxysporum f.sp. ciceri and half were left <sup>non</sup> inoculated. <sup>Non</sup> - Uncultivated beds were left for non-rhizosphere studies, Rhizosphere rhizoplane and non-rhizosphere studies were made from inoculated and uninoculated plants after 60 days.

Throughout the studies the petriplates were inoculated at 28°C unless stated otherwise. For each treatment there were 20 petriplates and each treatment was replicated thrice.

\* Experiments must be repeated at least one time. Indicate here that this was done to verify the results!!

describe how prepared.

# **Chapter Three**

## **Experimental Results**

## CHAPTER - III

### EXPERIMENTAL RESULTS

#### 3.1 RHIZOSPHERE AND RHIZOPLANE MYCOFLORA OF DIFFERENT CULTIVARS OF CHICKPEA INOCULATED WITH FUSARIUM OXYSPORUM F.SP. CICERI

*I will continue?*  
*There were*  
*species*  
*respectively*  
*respectively*

In all, 55 fungal species were isolated from the rhizosphere of six different cultivars of chickpea tested (Table 4). It was 28, 30, 29, 27, 28 and 27<sup>species</sup> in the rhizosphere of JG-62, H-208, BG-309, JG-315, BG-212 and JG-74<sup>respectively</sup> inoculated with Fusarium oxysporum f. sp. ciceri, as against 25, 26, 27, 23, 22 and 26<sup>respectively</sup> in the uninoculated plants. Thus in all the cultivars the inoculated plants harboured more fungi. In amongst inoculated ones highest number of fungi was recorded in H-208 and lowest in JG-74 and JG-315, while in uninoculated plants highest number of fungi was in BG-309 and lowest in BG-212. Aspergillus fumigatus, A. flavus, A. niger, Curvularia lunata, Drechslera hawaiiensis and Cladosporium herbarum were recorded from the rhizosphere of both inoculated and uninoculated plants of all the cultivars. However, Rhizopus oryzae, was isolated from all the cultivars except inoculated plants of JG-74. Rhizopus nodosus was recorded in the rhizosphere of H-208, BG-309, JG-315 and JG-74, Cunninghamella echinulata of JG-62, H-208, BG-309,

and BG-212; Monilia humicola of JG-62, JG-315, BG-212 and JG-74; Chaetomium flavum of H-208, BG-309, JG-315 and JG-62; Aspergillus funiculosus of H-208, BG-30 and BG-212; A. flavipes of JG-62 and H-208; A. clavatus of BG-309, BG-212 and JG-74; Botryotrichum piluliferum of BG-309, JG-315; Verticillium glaucum of JG-315 and BG-212; Memnoniella echinata of JG-309, BG-212 and JG-74; A. terreus of BG-309, JG-315 and JG-74; Curvularia geniculata of BG-212; Gliocladium roseum of JG-74; Penicillium notatum of H-208, BG-309 and JG-74; Acrophialophora fusipora of JG-62. Monosporium olivaceum, F. oxysporum f.sp. ciceri, Sclerotium rolfsii, Alternaria alternata and sterile brown mycelium were recorded from the rhizosphere of inoculated plants of all the cultivars whereas Mucor racemosus was confined to only JG-62, H-208, BG-309 and JG-315; Syncephalastrum racemosum of JG-62, H-208 and BG-212; Macrophomina phasiolina of JG-62, BG-309 and JG-315; Botrytis cinerea of JG-62, H-208 and BG-212; Nematogonium humicola of JG-315, Alternaria fasciculata to H-208, BG-309 and JG-315; Mortierella alpina to JG-62, H-208 and JG-74; and Sepdonium chrysospermum to JG-62, H-208, BG-309, JG-315; Rhizoctonia solani to JG-62, H-208 and BG-212. On the other hand, Memnoniella echinata, Trichoderma viridae and Humicola fuscotra were recorded from the rhizosphere of uninoculated plants. However, Phoma hibernica was isolated from all the cultivars except JG-62 whereas Aspergillus luchuensis was confined to JG-62, H-208

and JG-74, A. ochraceus to BG-212; A. sparsus to JG-74, H-208, and BG-309; Torulla allii to JG-62, H-208, BG-309 and JG-74; F. rosium to JG-31, BG-212; F. moniliformae to JG-62 and BG-309 and sterile black mycelium to JG-62, H-208, BG-309 and JG-74.

In amongst different fungi isolated from the rhizosphere of inoculated plants, the frequency of F. oxysporum f.sp. ciceri was highest in JG-62 followed by H-208, BG-309, BG-212, JG-315 and JG-74. A. fumigatus, A. flavus, A. niger, M. humicola, C. lunata and C. flavum in JG-315. M. racemosus, M. phaseolina A. flavipes, B. cineria, A. alternata in JG-62. S. chrysospermum C. herbarum and D. hawaiiensis in H-208. M. olivaceum, C. herbaru and D. hawaiiensis in BG-309. A. niger and D. hawaiiensis in JG-74. The frequency was lowest for A. flavus, A. niger, C. echinulata and C. lunata in JG-62; R. oryzae, A. fumigatus and sterile brown mycelium in H-20, R. nodosus, A. fumigatus in BG-309; M. olivaceum and A. alternata in BG-212 and A. niger and T. lignorum in JG-74. On the other hand, in rhizosphere of uninoculated plants the frequency of A. funigatus was highest in JG-315 and BG-212; A. flavus, C. lunata in JG-315; T. viridae in JG-315 and JG-74; C. aspermum, A. terreus, A. luchuensis, P. notatum, M. echinata and T. allii in JG-74. The frequency of C. echinulata, A. flavus, A. terreus, A. luchuensis was lowest in JG-62; R. oryzae, A. fumigatus, H. fuscotra, M. echinata, and C. lunata in H-208; F. oxysporum,

sterile black mycelium in BG-309; *C. flavum* in JG-315; *M. humicola* and *A. humicola* in BG-212.

In all 35 fungi were isolated from the rhizoplane of inoculated and uninoculated plants of six different cultivars of chickpea (Table-5). Thus the number of fungi was less in rhizoplane than in the rhizosphere. The number of fungi was more in the rhizoplane of inoculated plants of all the cultivars than in uninoculated plants. Highest number of fungi was found associated with the rhizoplane of H-208 and JG-62 in inoculated and uninoculated plants and lowest of BG-212 and JG-315 in uninoculated and inoculated plants. *Aspergillus fumigatus*, *A. flavus*, *A. niger*, *Cladosporium herbarum*, *Drechslera hawaiiensis* and *Curvularia lunata* were isolated from the rhizoplane of both inoculated and uninoculated plants of all the cultivars. However, *Cunninghamella echinulata* was reported from rhizoplane of JG-62, H-208, BG-309 and BG-212; *Rhizopus nodosus* of H-208, BG-309, JG-315 and JG-74; *R. oryzae* of JG-62, H-208, BG-309, JG-315 and BG-212; *Chaetomium flavum* of JG-62, H-208, BG-309 and JG-315; *Monilia humicola* of JG-315 and BG-212; *Gliocladium roseum* of JG-74; *Memnoniella echinata* of BG-309 and BG-212; *Aspergillus clavatus* of BG-212 and JG-74; *A. flavipes* of JG-62 and H-208 and *Acrophialophora fusipora* of JG-62. *Monosporium olivaceum*, *Alternaria alternata*, *Fusarium oxysporum* f.sp. *ciceri* and *Sclerotium rolfsii* were isolated from the rhizoplane of inoculated plants of all the different cultivars



whereas Mucor racemosus from all the cultivars except BG-212; Aspergillus candidus from H-208, BG-309 and BG-212, Mortierella alpina from JG-62, H-208 and JG-74; Botrytis cinerea from JG-62 and H-208; Sepdonium chrysospermum from H-208, BG-309, JG-315 and JG-74; Rhizoctonia solani from JG-62 and H-208. On the other hand A. terreus and sterile black mycelium were recorded from the rhizoplane of inoculated plants of all the cultivars whereas Torulla allii to H-208 and BG-309; Humicola fuscotra to JG-62, BG-212, JG-315 and JG-74; Aspergillus luchuensis to JG-62 and H-208.

The frequency of F. oxysporum f.sp. ciceri was highest in the rhizoplane of inoculated plants of all the cultivars except JG-315, BG-212 and JG-74 where the frequency of A. flavus and A. funigatus was highest. The frequency of A. flavus was highest in all the cultivars of uninoculated plants. On the other hand, the frequency was lowest for M. olivaceum in JG-62; R. oryzae, R. nodosus, M. racemosus and S. rolfsii in H-208; M. echinata in BG-309; R. nodosus, C. flavum and S. chrysospermum in JG-315; S. chrysospermum in BG-212 and R. nodosus, M. racemosus, M. alpina, A. alternata, S. rolfsii and T. viridae in JG-74 inoculated plants. In uninoculated plants the lowest frequency was that of C. echinulata in JG-62; A. terreus and A. flavipes in H-208, C. lunata in BG-309; and JG-315, M. humicola and C. herbarum in BG-212 and D. hawaiiensis and H. fuscotra in JG-74.

The population of fungi was highest in cv. H-208 both in inoculated and uninoculated plants (Table - 1). It was 185000 in inoculated as against 125000 in uninoculated plants thus indicating more population of fungi in rhizosphere of inoculated plants.

There <sup>was</sup> difference in the amino acid, phenol, O-dihydroxyphenole and sugar contents of different cultivars (Table-2).

The concentration of free amino acids was highest in cv. JG-74 both in inoculated and uninoculated plants. However sugar content was highest in cv. JG-62 in both inoculated and uninoculated plants. Amongst the two, the concentration was highest in inoculated plants root exudates.

Different cultivars of chickpea also exhibited qualitative differences in the amino acids (Table-3). The number of amino acids was highest in H-208, BG-309 and lowest in JG-74 and BG-212. Alanine, Aspartic acid Glutamic acid and Glycine were detected in the root extract of all the cultivars; Asparagine, Lysine and Histidine in JG-62, H-208, JG-309 and BG-212, Serine in H-208, BG-309, JG-315, BG-212 and JG-74; Tryptophane and Valine in JG-62, H-208 and JG-309, Tyrosine in JG-315, BG-212 and JG-74 and Lucine in JG-315. Phenylalanine in JG-62, H-208, BG-309, BG-212 and JG-74 and Cyteine in BG-212 and JG-74. The susceptible cultivars has higher number of free amino acids than resistant cultivars.

FIG. 1. Population of fungi in the rhizosphere of uninoculated and inoculated (with Fusarium oxysporum f.sp. ciceri) plants of different cultivars of chickpea.

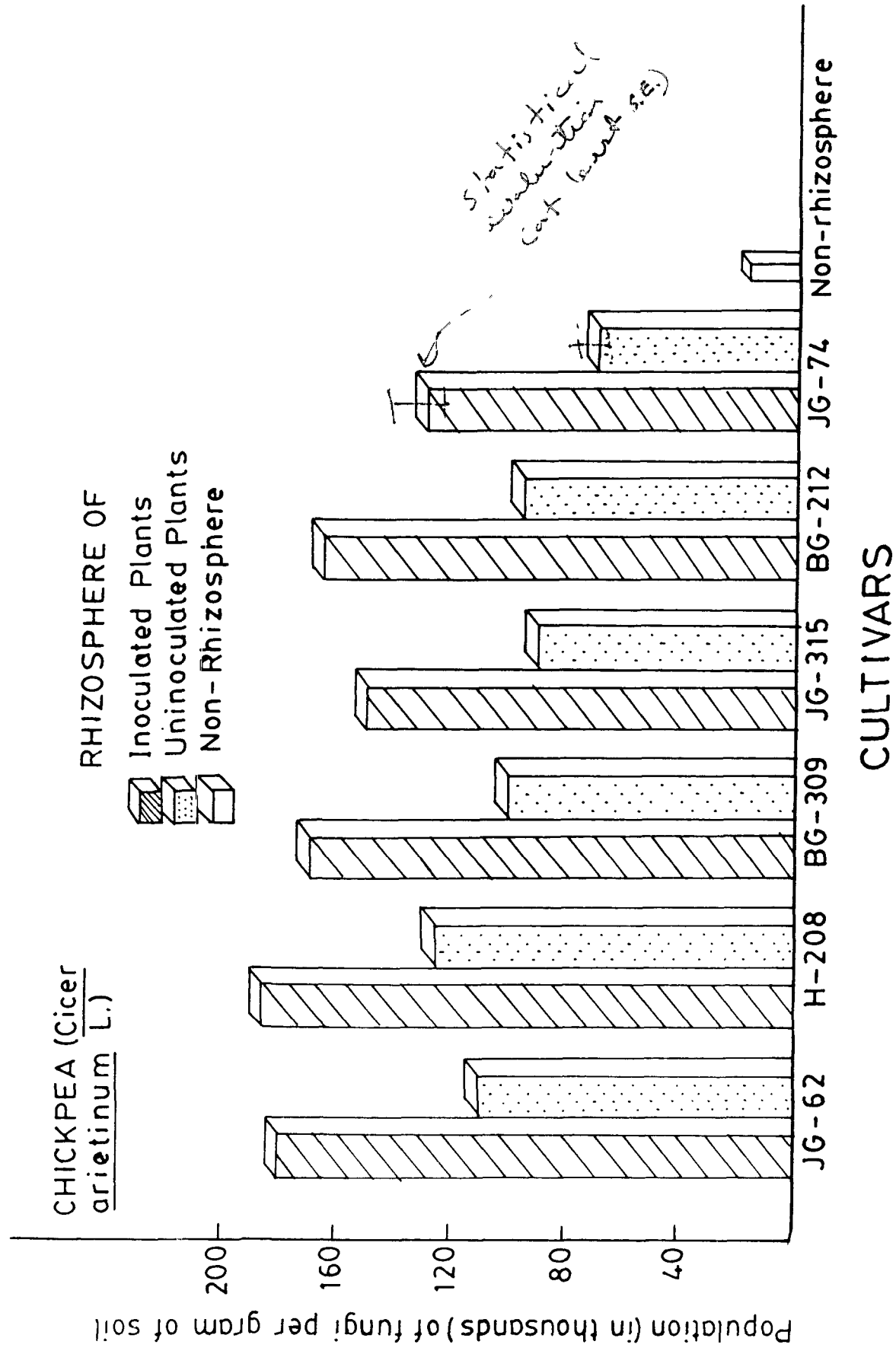


Fig.1

TABLE - 1: Population of fungi (per gram of soil) in different chickpea cultivars uninoculated and those inoculated with Fusarium oxysporum f.sp. ciceri

Caltivars	POPULATION			
	Inoculated Plants	R:S	Uninoculated Plants	R:S
Non-rhizosphere	17 <sup>+</sup>		17	
JG-62	180	10.58	110	6.47
H-208	185	10.88	125	7.35
BG-309	170	10.00	100	5.88
BG-212	150	8.82	90	5.29
JG-315	165	9.70	95	5.59
JG-74	130	7.64	70	4.11
L.S.D. at 5%	1.7888863		1.9651918	
L.S.D. at 1%	2.5444259		2.7951942	

<sup>+</sup>in thousands

Needs multiple comparisons to show signif. diff.

TABLE - 2: Concentrations (in mg/100 mg of dry samples) of free amino acids, phenols, O-dihydroxy phenols and sugars in the roots of different chickpea cultivars uninoculated and those inoculated with Fusarium oxysporum f.sp. ciceri

Cultivars	Free amino acids		Phenols		O-dihydroxy phenols		Sugars	
	IP	UP	IP	UP	IP	UP	IP	UP
JG-62	0.45	0.30	1.30	0.90	0.185	0.145	2.90	2.15
H-208	0.60	0.35	1.60	1.15	0.165	0.120	2.70	1.95
BG-309	0.70	0.50	2.70	1.90	0.280	0.200	2.50	1.75
JG-315	0.90	0.75	3.30	2.10	0.300	0.205	2.40	1.40
BG-212	1.05	0.80	3.60	2.50	0.375	0.265	1.35	1.05
JG-74	1.25	1.00	3.75	2.50	0.425	0.305	1.55	1.20
L.S.D. at 5%	0.095398	0.073500	0.098249	0.098244	0.0099638	0.0095309	0.098244	0.098244
L.S.D. at 1%	0.135690	0.104543	0.139752	0.139738	0.014172	0.0302034	0.139973	0.139973

IP = Inoculated plants

UP = Uninoculated plants

*Need multiple comparison*

TABLE - 3: Amino acids present in the roots of different chickpea cultivars uninoculated and those inoculated with Fusarium oxysporum f.sp. ciceri

Amino acids	CULTIVARS											
	JG-62		H-208		BG-309		JG-315		BG-212		JG-	
	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP
Alanine	+	+	+	+	+	+	+	+	+	+	+	+
Asparagine	+	+	+	+	+	+	+	+	-	-	-	-
Aspartic acid	+	+	+	+	+	+	+	+	+	+	+	+
Glutamic acid	+	+	+	+	+	+	+	+	+	+	+	+
Glycine	+	+	+	+	+	+	+	+	+	+	+	+
Histidine	+	+	+	+	+	+	+	+	-	-	-	-
Leucine	-	-	-	-	-	-	+	+	-	-	-	-
Lysine	+	+	+	+	+	+	+	+	-	-	-	-
Phenylalanine	+	+	+	+	+	+	-	-	+	+	+	+
Proline	+	+	+	+	+	+	-	-	-	-	-	-
Serine	-	-	+	+	+	+	+	+	+	+	+	+
Tryptophane	+	+	+	+	+	+	-	-	-	-	-	-
Tyrosine	-	-	-	-	-	-	+	+	+	+	+	+
Valine	+	+	+	+	+	+	-	-	-	-	-	-
Cysteine	-	-	-	-	-	-	-	-	+	+	+	+

IP = inoculated plants, UP = uninoculated plants.

TABLE - 4: Frequency (percentage) of fungi in the rhizosphere of different chickpea cultivars under IP and UP conditions. Those inoculated with Fusarium oxysporum f. sp. ciceri

Fungi Isolated	CULTIVARS											
	JG-62		H-208		BG-309		JG-315		BG-212		JG-24	
	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP
<u>Rhizopus oryzae</u>	20	30	10	20	20	25	15	25	10	30	-	25
<u>R. nodosus</u>	-	-	10	20	5	20	10	15	-	-	10	15
<u>Cunninghamella verticillatae</u>	-	-	-	-	20	-	20	10	15	-	-	-
<u>C. echinulata</u>	20	5	25	15	30	15	-	-	30	10	20	-
<u>Mucor racemosus</u>	25	-	15	-	20	-	15	-	-	-	-	-
<u>M. globosus</u>	-	-	-	-	-	-	-	-	15	-	20	10
<u>Mortierella alpina</u>	25	-	25	-	-	-	-	-	-	-	10	-
<u>Monilia humicola</u>	30	20	-	-	-	-	35	20	20	10	25	15
<u>Cephalosporium asperum</u>	10	20	15	25	-	-	-	20	-	-	-	25
<u>Syncephalastrum racemosum</u>	15	-	20	-	-	-	-	-	30	-	-	-
<u>Chaetomium flavum</u>	15	20	15	30	20	20	25	10	15	-	20	-
<u>Macrophomina phaseolina</u>	20	-	-	-	15	-	10	-	-	-	-	-
<u>Phoma hibemica</u>	-	-	-	25	-	20	-	15	-	20	-	10
<u>Aspergillus fumigatus</u>	35	25	25	20	25	20	45	45	40	35	10	30
<u>A. flavus</u>	20	25	35	40	25	40	60	65	55	65	25	35
<u>A. niger</u>	20	40	30	55	25	55	35	45	35	60	20	45
<u>A. terreus</u>	-	20	-	-	10	25	15	30	-	35	20	40
<u>A. luchuensis</u>	-	20	-	25	-	-	-	-	-	-	-	30
<u>A. funiculosus</u>	-	20	10	25	10	20	-	-	15	30	-	-
<u>A. candidus</u>	15	-	20	-	20	-	-	-	20	-	15	-
<u>A. flavipes</u>	30	20	20	10	-	-	-	-	-	-	25	-
<u>A. spargus</u>	10	25	-	30	-	20	-	-	-	-	-	20
<u>A. clavatus</u>	-	-	-	-	15	30	-	-	20	35	15	-
<u>A. ochraceus</u>	-	-	-	-	-	-	-	-	-	10	-	-
<u>Penicillium notatum</u>	-	-	10	20	15	30	-	-	-	25	20	35
<u>Gliocladium roseum</u>	-	-	-	-	-	-	-	-	-	-	25	35
<u>Monosporium olivaceum</u>	15	-	25	-	30	-	20	-	10	-	15	-
<u>Botrytis cinerea</u>	20	-	15	-	-	-	-	-	5	-	-	-
<u>Torula alii</u>	-	20	-	15	-	20	-	-	-	-	-	25
<u>Septonium chrysospermum</u>	15	-	30	-	20	-	20	-	15	-	-	-
<u>Botryotrichum pulluliferum</u>	-	-	-	-	20	30	20	15	-	-	-	-
<u>Verticillium glaucum</u>	-	-	10	-	-	-	15	5	20	10	25	-
<u>Cladosporium herbarum</u>	30	20	35	20	35	30	30	25	25	20	30	25
<u>Humicola fuscotra</u>	-	25	-	10	-	20	-	30	-	25	-	15
<u>Memnoniella echinata</u>	-	25	-	15	15	20	-	15	10	20	15	30
<u>Drechslera hawaiiensis</u>	35	30	40	35	40	30	30	25	40	30	40	30
<u>D. nodosus</u>	-	-	30	10	-	-	-	-	-	-	-	-
<u>Alternaria alternata</u>	30	15	25	-	15	-	20	-	10	-	20	-
<u>A. humicola</u>	-	-	-	20	-	15	-	20	15	10	-	20
<u>A. fusiculata</u>	-	-	20	-	20	-	20	-	-	-	-	-
<u>Curvularia lunata</u>	25	15	30	10	35	20	50	35	45	30	40	30
<u>C. geniculata</u>	-	-	-	-	-	-	-	25	15	-	-	-
<u>Fusarium roseum</u>	-	-	-	-	-	-	-	20	-	25	-	-
<u>F. moniliiformae</u>	-	20	-	-	-	35	-	-	-	-	-	-
<u>F. oxysporum</u>	-	35	-	25	15	20	15	25	-	-	-	-
<u>F. oxysporum</u> f.sp. <u>ciceri</u>	60	-	55	-	50	-	40	-	45	-	35	-
<u>Acremonialophora fusipora</u>	10	10	-	-	-	-	-	-	-	-	-	-
<u>Trichoderma viridae</u>	-	25	-	25	-	30	-	35	15	40	10	35
<u>T. lignorum</u>	-	-	-	-	-	20	15	20	-	15	10	15
<u>Hamatogonium humicola</u>	-	-	-	-	-	-	25	-	-	-	-	-
<u>Phloeoctonia solani</u>	20	-	15	-	-	-	-	5	-	-	-	-
<u>Sclerotium rolfsii</u>	15	-	10	-	20	-	10	-	5	-	15	-
Brown sterile mycelium	25	-	10	-	15	-	15	-	20	-	20	-
Black sterile mycelium	-	25	-	15	-	10	-	30	-	40	-	15
White sterile mycelium	10	-	15	15	10	10	25	-	-	-	-	-

IP - Inoculated plants; UP - Uninoculated plants.

S.E.   
 marked



TABLE - 5: Frequency (percentage) of fungi in the rhizoplane of different chickpea cultivars uninoculated and those inoculated with Fusarium oxysporum f.sp. ciceri

Fungi isolated	CULTIVARS											
	JG-62		H-208		BG-309		JG-315		BG-212		JG-74	
	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP
<u>Rhizopus oryzae</u>	25	35	10	30	15	25	15	30	20	30	-	30
<u>R. nodosus</u>	-	-	10	25	15	30	10	20	-	-	10	15
<u>Cunninghamella echinulata</u>	20	5	20	10	25	15	-	-	30	20	20	-
<u>Mucor racemosus</u>	20	-	10	-	15	-	15	-	-	-	10	-
<u>Mortierella alpina</u>	20	-	15	-	-	-	-	-	-	-	10	-
<u>Monilia humicola</u>	-	-	-	-	-	-	30	20	25	10	-	-
<u>Cephalosporium asperum</u>	-	30	-	25	-	-	-	30	-	-	-	-
<u>Chaetomium flavum</u>	15	25	15	30	20	25	10	20	-	-	-	-
<u>Aspergillus fumigatus</u>	30	20	40	30	25	25	40	30	50	40	40	30
<u>A. flavus</u>	25	40	40	50	25	40	50	60	40	70	30	45
<u>A. niger</u>	20	35	30	50	15	40	25	60	35	70	35	40
<u>A. terreus</u>	-	20	-	10	-	30	-	25	-	25	-	30
<u>A. luchuensis</u>	-	20	-	30	-	-	-	-	-	-	-	-
<u>A. candidus</u>	-	-	30	-	35	-	-	-	25	-	-	-
<u>A. clavatus</u>	-	-	-	-	-	-	-	-	20	30	20	35
<u>A. sparsus</u>	15	20	-	25	-	20	-	-	-	-	-	20
<u>A. flavipes</u>	25	15	30	10	-	-	-	-	-	-	-	-
<u>Gliocladium roseum</u>	-	-	-	-	-	-	-	-	-	-	35	25
<u>Monosporium olivaceum</u>	10	-	40	-	35	-	20	-	20	-	15	-
<u>Botrytis cinerea</u>	20	-	15	-	-	-	-	-	-	-	-	-
<u>Cladosporium herbarum</u>	20	15	25	15	30	20	25	15	15	10	30	20
<u>Memnoniella echinata</u>	-	30	-	15	10	20	-	15	15	20	-	25
<u>Drechslera hawaiiensis</u>	20	15	40	20	35	25	40	25	45	35	25	15
<u>Curvularia lunata</u>	35	25	30	15	20	10	25	10	40	20	35	25
<u>Alternaria alternata</u>	20	10	15	-	15	-	20	-	15	-	10	-
<u>Fusarium oxysporum f.sp. ciceri</u>	60	-	60	-	55	-	50	-	45	-	40	-
<u>Acrophialophora fusipora</u>	15	10	-	-	-	-	-	-	-	-	-	-
<u>Trichoderma viridae</u>	-	25	-	20	-	25	-	35	10	40	10	35
<u>Torula alli</u>	-	-	-	15	-	25	-	-	-	-	-	-
<u>Scedonum chrysospermum</u>	-	-	-	-	15	-	10	-	5	-	-	-
<u>Humicola fuscoatra</u>	-	25	-	-	-	-	-	15	-	20	-	15
<u>Rhizoctonia solani</u>	20	-	15	-	-	-	-	-	-	-	-	-
<u>Sclerotium rolfsii</u>	15	-	10	-	20	-	15	-	15	-	10	-
<u>Sterile brown mycelium</u>	20	-	20	-	-	-	25	-	-	-	25	-
<u>Sterile black mycelium</u>	-	20	-	30	-	30	-	25	-	45	-	30

Added 5 strains  
analysis

IP = Inoculated plants; UP = Uninoculated plants

### 3.2 EFFECT OF AGE OF CHICKPEA CULTIVAR JG-62 ON THE RHIZOSPHERE MYCOFLORA INOCULATED WITH FUSARIUM OXYSPORUM F.SP. CICERI

Out of 45 fungal species isolated, the number of fungi from rhizosphere of inoculated plants was 34 as against 24 and 30 from non-rhizosphere zone and the rhizosphere of uninoculated plants respectively (Table-8). As a result of inoculation, differences were observed in the rhizosphere mycoflora Mucor racemosus, Cunninghamella echinulata, Mortierella alpina, Macrophomina phaseolina, Syncephalastrum racemosum, Aspergillus candidus, Botrytis cinerea, Monosporium olivaceum, Rhizoctonia solani and sterile brown mycelium were confined to the rhizosphere of inoculated plants in addition to Fusarium oxysporum f.sp. ciceri with which the plants were inoculated and Aspergillus terreus, Botryotrichum piluliferum and sterile black mycelium in the rhizosphere of uninoculated plants whereas Geotrichum candidum, Aspergillus koningi and sterile white mycelium in the non-rhizosphere area.

The number of fungi in the rhizosphere of both inoculated and uninoculated plants increased with the increase in the age of plants upto 60 days followed by a decline with more fungi in inoculated plants (Table-8). Aspergillus fumigatus, A. flavus, A. niger, Monilia humicola and Cladosporium herbarum were recorded throughout the growth period in the rhizosphere of

both inoculated and uninoculated plants. Aspergillus funiculosus, Fusarium oxysporum and Trichoderma viridae were isolated in the early part of growth period, Aspergillus sparsus, Rhizopus oryzae, Acrophialophora fusipora and Alternaria alternata from early to intermediate growth period; Cephalosporium aspermum, Aspergillus okazakii, A. flavipes and Trichosporium fuscum during intermediate growth stage; Chaetomium flavum, Drechslera hawaiiensis, Curvularia lunata from intermediate to later part of growth period and A. funiculosus and Spicaria silvatica in the later part of growth period. from rhizosphere of both inoculated and uninoculated plants. In the rhizosphere of inoculated plants Mucor racemosus, C. echinulata, A. flavipes, Mortierilla alpina F. oxysporum f.sp. ciceri, A. alternata and sterile brown mycelium were recorded throughout the growth period; P. lilacinum in the early growth period; A. candidus, S. racemosum from early to intermediate growth stages; B. cinerea during intermediate growth stage; M. phasiolina, S. rolfsii and S. chrysospermum from intermediate to later part of the growth period and R. solani in later part of growth period, while from the rhizosphere of uninoculated plants A. luchuensis, T. allii, M. echinata and sterile black mycelium at all the intervals of growth period; D. piluliferum in the early part; Fusarium moniliformae during intermediate and A. terreus, and Humicola fuscotra from intermediate to later part of growth period.

There was increase in frequency of most of the fungi in the rhizosphere of both inoculated and uninoculated plants with the increase in age upto a certain period followed by a decline. In some cases there was again an increase. However, in the non-rhizosphere no definite pattern of frequency of different fungi was observed. In the rhizosphere of inoculated plants frequency of certain fungi exhibited a decline with increase in age of plants. These fungi include R. oryzae, A. flavus, A. niger and D. hawaiiensis. Frequency of F. oxysporum f.sp. ciceri in the rhizosphere of inoculated plants increased with the increase in age of plant upto 75 days then slightly decreased with onset of senescence. In the rhizosphere of inoculated plants, the frequency of most of the saprophytic fungi decreased with the increase in age of plant, however the reverse was true with the parasitic fungi. The frequency of saprophytic fungi was higher in the rhizosphere of uninoculated plants while that of parasitic fungi in the rhizosphere of inoculated plants.

The number of fungi on the rhizoplane of chickpea plants inoculated and uninoculated was slightly less than corresponding rhizosphere and it was 27 and 24 fungal species respectively (Table-9). The number of fungi in the rhizoplane also increased with the increase in age of plant upto a stage after which there was a decrease. Not all fungi were isolated from rhizoplane at

all the stages of plant growth. Aspergillus funigatus, A. flavus, A. niger and Cladosporium harbarum were recorded in the rhizoplane of both inoculated and uninoculated plants at all the stages of the growth of plant. However, other fungi were present at certain stages of plant growth only. Mucor racemosus, Cunninghamella echinulata, Aspergillus funiculus, Alternaria alternata and Acrophialophora fusipora were isolated from early part of growth of plant Rhizopus oryzae, Aspergillus sparsus from early to middle age Sclerotium rolfsii in the middle age; Curvularia lunata, Drechslera hawaiiensis and Aspergillus ochroaceus from middle to last stages of plant growth in the rhizoplane of both inoculated and uninoculated plants. However, in the rhizoplane of inoculated plants Monosporium olivaceum, Macrophomina phasiolina and Fusarium oxysporum f.sp. ciceri at all stages of plant growth; Botrytis cinerea, Mucor racemosus in the middle age of growth of plant and Alternaria alternata, Rhizoctonia solani and Sclerotium rolfsii in the last phases of growth of plant. In the rhizoplane of uninoculated plants Memnoniella echinata, Aspergillus luchuensis, sterile black mycelium were recorded at all the stages of growth of plant; Aspergillus terreus, Humicola fuscotra, Trichoderma viridae and Cephalosporium aspermum from middle to last phases of growth of plant and R. oryzae and A. funiculosus. In the later stages of plant growth, M. alpina, M. phasiolina, B. cinerea, M. olivaceum, R. solani

and sterile brown mycelium were exclusively present in the rhizoplane of inoculated plants and A. luchuensis, A. terreus, Fusarium oxysporum, M. echinata, Cephalosporium aspermum, H. fuscotra and sterile black mycelium in that of uninoculated plants only.

For most of the fungi, the frequency increased with increase in age of plants upto 60 days followed by a decline. However, in the rhizoplane of uninoculated plants the frequency of A. flavus increased upto senescence. In the rhizoplane of inoculated plants certain fungi such as A. fumigatus, A. flavus and A. niger exhibited a decline with increase in age of plants. The frequency of F. oxysporum f.sp. ciceri in the rhizoplane of inoculated plants increased with increase in age of plants upto 75 and then remain constant. In the rhizoplane of inoculated plants, the frequency of most of the saprophytic forms decreased with the increase in age of the plant however, the reverse was true with the parasitic fungi.

The population of fungi in the rhizosphere of inoculated and uninoculated plants also increased in age of plant upto 75 days followed by decline (Table-6). In inoculated plants, however, there was again an increase at 120-days-old plants. The population of fungi was high in the rhizosphere of inoculated plants as compared to uninoculated ones.

When compared with non rhizosphere mycoflora the R:S ratio increased with increase in the age of plant upto 90 days in inoculated plants as against 75 days in uninoculated plants (Table-6). However, highest R:S ratio in inoculated plants was at 120 days old. Thus the inoculated plants provide more favourable environment for the rhizosphere fungi.

Quantitative analysis of the root extracts indicated that the concentration of free amino acids (Mg/100 mg of dry sample) varied at different stages of plant growth (Table- 7). It also exhibited an increase upto certain stage of plant growth followed by decline in both root extracts of inoculated and uninoculated plants. A similar trend was observed with total phenols, O-dihydroxyphenols and sugar contents in the roots' extracts of both inoculated and uninoculated plants. The concentration of free amino acids, phenols, O-dihydroxyphenols and sugars was high in the roots of inoculated plants as compared to uninoculated ones.

FIG. 2. Population of fungi in the non-rhizosphere, rhizosphere of uninoculated and inoculated plants of chickpea (with Fusarium oxysporum f.sp. ciceri) at varying age of the plant.



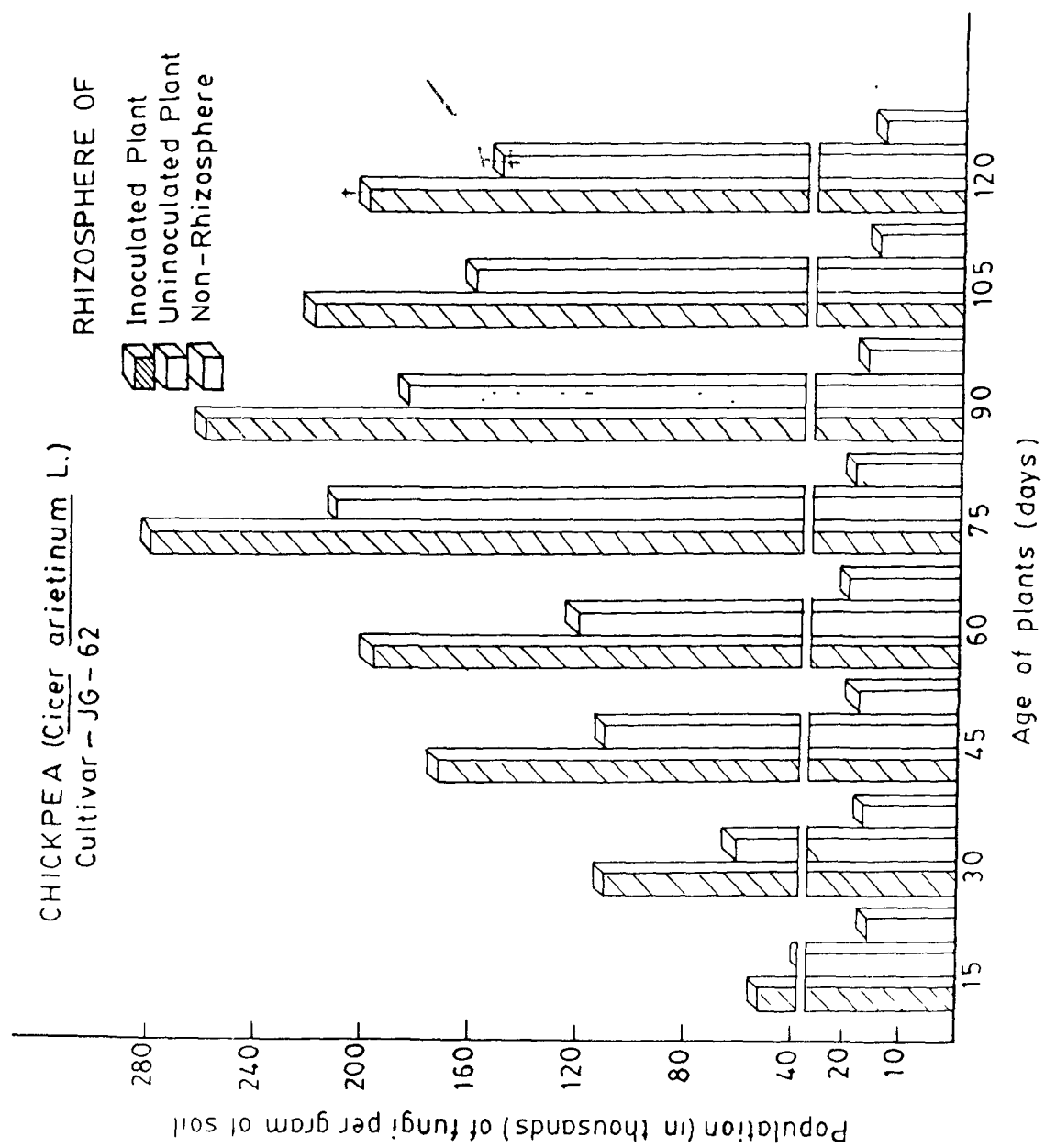


Fig.2

TABLE - 6: Population of fungi (per gram of soil) in non-rhizosphere, rhizosphere of uninoculated and those inoculated with Fusarium oxysporum f.sp. ciceri of chickpea plant cultivar JG-62 at varying age

Age of plants (days)	Non-rhizosphere(S)	RHIZOSPHERE (R)		
		Inoculated Plants	R:S Ratio	Uninoculated Plants R:S Ratio
15	16 <sup>+</sup>	51	3.18	34 2.12
30	17	110	4.23	61 3.58
45	18	172	6.11	110 4.61
60	20	196	9.80	120 6.00
75	19	280	14.73	210 11.05
90	17	260	15.29	185 10.88
105	15	220	13.33	160 10.66
120	14	200	15.71	150 10.71
L.S.D. at 5%	1.7669531	1.7038271		1.8576244
L.S.D. at 1%	3.2760712	3.1590305		3.444183

<sup>+</sup> in thousands

Needs statistical comparison (multivariately)  
~~statistical~~ comparison?   
 linear perhaps?

TABLE - 7: Concentration (in mg/100 mg of dry sample) of free amino acids, phenols, O-dihydroxy-phenols, sugars in the roots of chickpea plant cultivar JG-62 uninoculated and those inoculated with Fusarium oxysporum f.sp. ciceri at varying age

Age of Plant	Free amino acids		Phenols		O-dihydroxyphenols		Sugars	
	IP	UP	IP	UP	IP	UP	IP	UP
30	0.28	0.19	0.75	0.40	0.10	0.08	2.45	1.90
45	0.40	0.25	1.15	0.85	0.175	0.125	2.75	2.20
60	0.76	0.45	1.70	1.35	0.295	0.225	3.55	2.60
75	0.95	0.62	1.85	1.55	0.325	0.245	3.65	2.85
90	0.82	0.70	1.65	1.20	0.250	0.145	3.30	2.50
105	0.76	0.60	1.25	1.05	0.175	0.110	3.00	2.10
120	0.64	0.50	0.95	0.95	0.135	0.100	2.80	1.95
L.S.D. at 5%	0.0316297	0.0346473	0.0228919	0.0216613	0.0023347	0.0020729	0.847202	0.068303
L.S.D. at 1%	0.0435804	0.0477381	0.0315412	0.0298456	0.0032168	0.0028561	0.1167301	0.094109

TABLE - 8 Frequency of isolation of fungi in the rhizosphere of *Phaseolus* plant (cultivar SG-62) uninoculated and those inoculated with *Aspergillus* f.sp. *ciceri* at varying age

Fungi isolated	15			30			45			60			75			90			105			120		
	R		NR	R		NR	R		NR	R		NR	R		NR	R		NR	R		NR	R		NR
	IP	UP		IP	UP		IP	UP		IP	UP		IP	UP		IP	UP		IP	UP		IP	UP	
<i>Rhizopus oryzae</i>	30	30	35	25	30	25	20	30	35	10	30	30	10	20	25	15	15	15	15	15	15	15	15	15
<i>Mucor racemosus</i>	0	-	-	15	-	-	15	-	-	0	-	-	10	-	-	10	-	-	15	-	-	-	-	-
<i>Cunninghamella echinulata</i>	20	-	15	25	-	20	30	-	20	25	20	-	15	10	-	10	-	-	-	-	-	10	-	-
<i>Mortierella alpina</i>	10	-	10	10	-	10	15	-	15	5	-	10	20	-	10	10	-	-	-	-	-	-	-	-
<i>Geotrichum candidum</i>	-	-	20	-	-	35	-	-	35	-	-	35	-	-	30	-	-	30	-	-	20	-	-	25
<i>Monilia humicola</i>	15	10	-	20	15	-	30	25	-	40	30	-	35	20	-	25	15	-	15	10	-	10	5	-
<i>Cephalosporium as</i>	-	-	-	15	10	-	10	15	-	5	25	-	-	20	-	-	10	-	-	-	-	-	-	-
<i>Chaetomium flavum</i>	-	-	-	-	-	-	20	20	-	20	25	15	35	20	10	40	15	-	40	10	-	40	10	10
<i>Macrophoma phaseoli</i>	-	-	-	15	-	-	20	-	-	20	-	-	25	-	-	25	-	-	20	-	-	15	-	-
<i>Aspergillus fumigatus</i>	40	40	20	40	30	25	40	25	20	40	30	20	40	30	20	35	30	20	30	20	25	20	25	15
<i>A. flavus</i>	40	20	10	35	20	10	25	30	15	30	50	10	20	0	20	10	60	25	20	70	35	20	75	30
<i>A. niger</i>	15	20	25	20	30	20	20	35	25	40	55	20	30	40	25	30	35	30	25	20	30	20	20	20
<i>A. funiculosus</i>	1	-	10	20	-	-	15	-	-	15	-	-	15	-	-	20	10	-	-	-	10	-	20	10
<i>A. terreus</i>	-	-	-	-	-	-	20	-	-	20	-	-	20	-	-	20	-	-	1	-	-	10	-	-
<i>A. luchuensis</i>	-	1	10	-	10	-	-	20	10	-	0	-	-	25	-	-	15	-	-	10	-	-	10	-
<i>A. candidus</i>	10	-	-	10	-	-	15	-	-	-	-	-	15	-	-	5	-	-	-	-	-	-	-	-
<i>A. okazaki</i>	-	-	-	-	-	-	-	-	-	-	-	-	5	30	-	10	30	-	-	2	-	-	20	-
<i>A. flavipes</i>	-	-	-	25	-	-	30	10	-	25	20	-	25	15	-	30	-	-	35	-	-	30	-	-
<i>A. parvus</i>	0	15	10	20	25	-	20	25	10	5	30	15	10	40	20	-	25	-	-	-	-	10	-	-
<i>A. coningii</i>	-	-	10	-	-	15	-	-	20	-	-	25	-	-	20	-	-	25	-	-	20	-	-	15
<i>Penicillium lili</i>	-	-	-	15	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
<i>Botrytis cinerea</i>	-	-	-	-	-	-	15	-	-	20	-	-	20	-	-	10	-	-	-	-	-	-	-	-
<i>Torula alli</i>	-	15	-	-	15	-	-	20	10	-	25	-	-	20	20	-	25	20	-	15	15	-	10	10
<i>Monosporium olivaceum</i>	10	-	-	10	-	-	15	-	-	20	-	-	20	-	-	15	-	-	15	-	-	15	-	-
<i>Sepdonium chrysospermum</i>	-	-	-	-	-	-	-	-	-	15	-	-	20	-	-	30	-	20	20	-	20	25	-	10
<i>Botryotrichum pillularum</i>	-	15	-	-	10	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
<i>Cladosporium verburum</i>	20	10	-	20	20	-	30	25	-	40	30	-	35	30	-	30	20	-	30	25	-	20	10	-
<i>Spicaria silvatica</i>	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	30	15	-	25	20	-	20	20	-
<i>Syncephalastrum racemosum</i>	10	-	-	20	-	-	25	-	-	35	-	-	30	-	-	15	-	-	-	-	-	-	-	-
<i>Trichosporium fuscum</i>	-	-	-	-	-	-	-	-	-	30	20	-	20	15	-	15	10	-	-	-	-	-	-	-
<i>Humicola fuscora</i>	-	-	-	-	-	-	-	15	-	-	20	-	-	25	10	-	20	20	-	10	5	-	15	5
<i>Memnoniella ecinata</i>	-	10	-	-	15	-	-	20	-	-	25	10	-	25	15	-	20	15	-	20	10	-	20	-
<i>Drechslera hawaiiensis</i>	-	-	-	-	15	35	30	20	35	30	20	30	20	15	25	15	10	20	10	10	20	10	10	10
<i>Curvularia lunata</i>	-	-	-	-	-	-	20	20	-	25	20	-	30	25	-	25	10	-	25	10	-	30	10	-
<i>Alternaria alternata</i>	2	10	15	20	10	20	25	20	-	30	20	-	35	10	-	15	-	-	10	-	-	-	-	-
<i>Fusarium moniliforme</i>	-	-	-	-	-	-	20	-	-	25	-	-	15	-	-	15	-	-	-	-	-	-	-	-
<i>F. oxysporum</i>	2	25	15	10	25	20	-	25	-	15	-	-	-	-	-	-	-	-	-	-	-	-	-	-
<i>F. oxysporum f. sp. ciceri</i>	5	-	-	45	-	-	50	-	-	65	-	-	90	-	-	70	-	-	60	-	-	60	-	-
<i>Acronialophora f. ciceri</i>	15	10	10	20	10	15	25	15	-	30	15	-	15	-	-	-	-	-	-	-	-	-	-	-
<i>Trichoderma viride</i>	10	25	-	-	30	-	-	30	-	-	35	-	-	30	-	-	25	-	-	25	-	-	25	-
<i>Rhizoctonia solani</i>	-	-	-	-	-	-	15	-	10	20	-	-	-	-	-	15	-	-	10	-	-	10	-	-
<i>Sclerotium oryzae</i>	-	-	-	-	-	-	10	-	-	15	10	-	20	-	-	10	-	-	10	-	-	10	-	-
White triangle	-	10	-	-	15	-	-	20	-	-	15	-	-	15	-	-	10	-	-	10	-	-	1	-
Stirring brown	10	-	-	15	-	-	25	-	-	20	-	-	10	-	-	10	-	-	15	-	-	20	-	-
sterile leaf	10	-	-	15	-	-	20	-	-	30	-	-	25	-	-	20	-	-	25	-	-	-	-	-

R = radius  
IP = Inoculated plants

Statistical  
(S.E. not known)

TABLE - 9: Frequency (percentage) of fungi in the rhizoplane of chickpea plants (cultivar JG-62) uninoculated and those inoculated with Fusarium oxysporum f.sp. ciceri at varying age

	DAYS															
	15		30		45		60		75		90		105		120	
	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP
<u>Rhizopus oryzae</u>	25	15	30	25	35	30	25	40	15	45	-	35	-	25	-	15
<u>Gunninghamella echinulata</u>	15	10	20	5	25	-	30	-	20	-	10	-	15	-	10	-
<u>Mucor racemosus</u>	10	10	20	10	25	-	25	-	20	-	10	-	-	-	-	-
<u>Mortierella alpina</u>	10	-	15	-	20	-	20	-	15	-	10	-	-	-	-	-
<u>Macrophomina phaseolina</u>	-	-	10	-	15	-	20	-	20	-	15	-	15	-	10	-
<u>Chaetomium flavum</u>	-	-	10	15	20	20	20	25	30	25	30	10	25	10	15	10
<u>Aspergillus fumigatus</u>	45	40	45	40	35	25	50	30	40	30	40	30	30	25	20	20
<u>A. flavus</u>	35	35	30	40	25	40	25	55	20	60	20	65	25	65	25	70
<u>A. niger</u>	20	25	25	30	20	30	20	35	20	40	15	30	10	25	10	20
<u>A. terreus</u>	-	-	-	-	-	15	-	25	-	20	-	20	-	15	-	10
<u>A. luchuensis</u>	-	10	-	15	-	20	-	25	-	15	-	10	-	10	-	20
<u>A. okazaki</u>	-	-	-	-	-	-	-	-	15	30	10	25	-	20	-	10
<u>A. funiculosus</u>	20	20	15	10	-	-	-	-	-	15	-	10	-	15	-	25
<u>A. flavipes</u>	10	-	15	-	20	15	30	-	25	-	30	-	30	-	35	-
<u>A. ochraceus</u>	-	-	-	-	-	-	-	-	15	20	20	30	25	45	20	35
<u>A. sparus</u>	10	15	15	20	15	25	20	30	10	15	-	5	-	-	-	-
<u>Botrytis cinerea</u>	-	-	-	-	15	-	20	-	10	-	-	-	-	-	-	-
<u>Memnoniella echinata</u>	-	10	-	10	-	20	-	30	-	25	-	15	-	15	-	10
<u>Gladiosporium herbarum</u>	15	15	15	10	30	15	35	20	40	15	45	15	30	20	30	25
<u>Drechlera hawaiiensis</u>	-	-	10	-	15	10	20	15	30	15	30	10	30	10	25	-
<u>Curvularia lunata</u>	-	-	15	-	15	10	20	15	30	25	40	30	40	35	40	40
<u>Humicola fuscoatra</u>	-	-	-	-	-	10	-	15	-	15	-	10	-	5	-	-
<u>Alternaria alternata</u>	10	10	20	15	25	10	30	-	15	-	10	-	10	-	-	-
<u>Trichoderma viridae</u>	10	20	10	20	-	25	-	30	-	35	-	35	-	30	-	25
<u>Fusarium oxysporum</u>	-	25	-	15	-	10	-	-	-	-	-	-	-	-	-	-
<u>Fusarium oxysporum f.ciceri</u>	25	-	35	-	55	-	70	-	75	-	60	-	60	-	60	-
<u>Monosporium olivaceum</u>	10	-	10	-	25	-	30	-	30	-	25	-	20	-	15	-
<u>Cephaesporium aspermum</u>	-	-	-	15	-	20	-	30	-	25	-	25	-	10	-	10
<u>Acrophialophora fusipora</u>	10	10	15	10	20	15	-	10	-	-	-	-	-	-	-	-
<u>Rhizoctonia solani</u>	-	-	-	-	10	-	-	-	-	-	-	-	15	-	20	-
<u>Sclerotium rolfsii</u>	-	-	-	-	15	10	15	15	-	-	-	-	10	-	10	-
Sterile brown mycelium	10	-	15	-	25	-	30	-	20	-	15	-	20	-	15	-
Sterile black mycelium	-	15	-	25	-	30	-	20	-	10	-	-	-	-	-	-

IP = Inoculated plants, UP = Uninoculated plants.

Statistics  
(S.E. & C.V.)

3.3 RHIZOSPHERE AND RHIZOPLANE MYCOFLORA OF CHICKPEA PLANTS  
(CULTIVAR JG-62 AND JG-74) INOCULATED WITH FUSARIUM  
OXYSPORUM F.SP.CICERI IN RELATION TO FOLIARY SPRAY

(a) Growth Regulator:

Results presented in Tables 12 and 16 show that as a result of spray of plants with growth promoting substances, the number of fungi in the rhizosphere increased with the increase the number of sprays. The number of fungi in the rhizosphere was also more in those inoculated with F. oxysporum f.sp. ciceri. In amongst the two varieties tested the highest number of fungi was reported from the rhizosphere of JG-62 both in inoculated and those uninoculated. When sprayed with indole acetic acid, the number of fungi in JG-62 was 20, 26, 29 in inoculated as against 19, 25, 27 in uninoculated after respective sprays; with indole butyric acid 21, 26, 29 in inoculated as against 21, 24, 27 in uninoculated with thio indole butyric acid 19, 27, 29 in inoculated as against 19, 26, 28 in uninoculated; with gibberellic acid 19, 26, 29 in inoculated as against 20, 24, 27 in uninoculated, and with maleic hydrazide 18, 24, 26 in inoculated as against 17, 23, 26 in uninoculated plants in respective sprays. The number of fungi as a result of spray with water as a control was 19, 26, 29 in inoculated as against 19, 24, 27 in uninoculated plants. With JG-74, the corresponding values were 22, 28, 28 in inoculated as against

22, 26, 27 in uninoculated when sprayed with indole acetic acid 21, 27, 28 as against 21, 25, 23 with indole butyric acid, 21, 28, 28 as against 21, 26, 27 with thio-indole butyric acid, 22, 28, 28 as against 21, 26, 27 with gibberellic acid, and 21, 26, 25 as against 18, 25, 27 with maleic hydrazide while the number of fungi in the rhizosphere of plants sprayed with water as control was 21, 26, 28 as against 19, 25, 27 in uninoculated plants. A similar trend was observed in the increase in the number of fungi with number of sprays was obtained in the rhizoplane with more number of fungi in those inoculated with F. oxysporum f.sp. ciceri (Tables, 14 and 18). In amongst the two cultivars tested, number of fungi was more in the rhizoplane of the JG-62 both in inoculated and uninoculated plants. When sprayed with indole acetic acid, the number of fungi was 17, 20, 23 in the rhizoplane of inoculated plants of JG-62 as against 16, 19, 20 in respective sprays in uninoculated, with indole butyric acid 17, 21, 23 in inoculated as against 15, 19, 20 in uninoculated plants; with thio indole butyric acid 17, 20, 23 in inoculated as against 15, 19, 20 in uninoculated plants, with gibberellic acid 17, 20, 23 in inoculated as against 15, 19, 20 in uninoculated plants, and with maleic hydrazide 15, 18, 21 in inoculated as against 15, 19, 17 in uninoculated with respective sprays. With water as control, the number was 15, 20, 23 in inoculated as against 15, 19, 20 in uninoculated plants. With cv. JG-74 the corres-

ponding values were 15, 19, 20 as against 14, 18, 19 when sprayed with indole butyric acid; 15, 19, 20 as against 14, 18, 19 with indole butyric acid 15, 19, 20 as against 14, 18, 19 with thio-indole butyric acid; 15, 19, 20 as against 14, 18, 19 with gibberellic acid; and 14, 17, 19 as against 14, 18, 18 with maleic hydrazide. The number of fungi was 15, 18, 20 as against 14, 18, 19 after respective sprays with water. The number of fungi was less in rhizoplane as compared to rhizosphere. It appears there has not been material effect on the number of fungi both in rhizosphere and rhizoplane as a result of sprays with different growth promoting substances. By and large the population of fungi in the rhizosphere increased due to sprays with growth promoting substances and highest increase was observed when sprayed with indole acetic acid in inoculated plants (Table 10 and 11). In uninoculated as compared to inoculated ones there was less increase when compared with plants sprayed with water.

In general there was increase in the frequency as a result of sprays with growth promoting substances in the rhizosphere of JG-62 inoculated plants (Table 12 and 16) with spray of indole acetic acid highest increase in frequency of fungi was that of Macrophomina phaseolina followed by Monosporium olivaceum, Cladosporium herbarum, Fusarium oxysporium f.sp. ciceri, Aspergillus flavipes and Alternaria



alternata. However, in uninoculated plants the highest increase in the frequency was that of Cephalosporium asperum, followed by Aspergillus niger, Spicaria silvatica, Rhizopus oryzae and Monilia humicola with indole butyric acid highest increase in frequency of fungi was that of M. phaseolina followed by F. oxysporum f.sp. ciceri, Sclerotium rolfsii R. oryzae, Mortierilla alpina, Syncephalastrum racemosum, Aspergillus fumigatus, A. flavus in inoculated and that of Chaetomium flavum followed by Curvularia lunata, Fusarium moniliiformae and sterile black mycelium in uninoculated plants with thio indole butyric acid, R. oryzae followed by A. alternata, F. oxysporum f.sp. ciceri, sterile brown mycelium Rhizoctonia solani, C. lunata and M. olivacium in inoculated that of C. flavum followed by Aspergillus sparsus, A. flavipes, Humicola fuscotra, Curvularia geniculata and R. oryzae in uninoculated plants; with gibberllic acid F. oxysporum f.sp. ciceri followed by Mucor racemosus, S. rolfsii, A. flavipes, C. lunata, C. herbarum, S. racemosum and sterile brown mycelium in inoculated that of R. oryzae followed by A. fumigatus, A. flavus, A. terreus, A. luchuensis, C. herbarum, H. fuscotra, Memnoniella echinata, Drechslera hawaiiensis and C. lunata in uninoculated plants; with maleic hydrazide the frequency of most of the fungi decreased, the frequency of M. racemosus decreased highest followed by C. herbarum, Aspergillus sparsus A. flavipes, A. candidus, C. asperum, C. flavum, A. alternata,

and F. oxysporum f. ciceri inoculated that of A. fumigatus followed by A. flavipes, A. flavus, Acrophialophora fusipora in uninoculated plants, with an other variety JG-74 highest increase in the frequency was that of Mortierilla alpina followed by A. flavus, A. candidus, Penicillium notatum, S. rolfsii, T. viridae and sterile brown mycelium when sprayed with indole acetic in inoculated that of C. aspermum followed by Gliocladium roseum, Phoma hibernica, C. herparum and Trichoderma lignorum in uninoculated plants; with indole butyric acid highest increase in the frequency was that of Aspergillus clavatus followed by A. flavus, A. flavipes sterile brown mycelium and Mortierilla alpina in inoculated that of A. humicola followed by A. fumigatus, A. niger, T. viridae, H. fuscotra and sterile black mycelium in uninoculated plants; with thio indole butyric acid R. nodosus followed by A. flavus, A. fumigatus, A. niger, M. olivacium A. alternata, S. rolfsii and sterile brown mycelium in inoculated that of A. fumigatus followed by A. flavus, P. notatum, G. roseum H. fuscotra, P. hibernica and T. viridae and sterile black mycelium in uninoculated plants; with gibberellic acid M. globosus followed by A. candidus, A. flavus, A. fumigatus, A. alternata, T. viridae, S. rolfsii and sterile brown mycelium in inoculated that of P. hibernica followed by A. clavatus, A. terreus, H. fuscotra, M. echinata, C. lunata, A. humicola and sterile black mycelium in

uninoculated plants. But with maleic hydrazide the frequency of M. racemosus, A. terreus and C. herbarum decreased most in inoculated that of P. hibernica, H. fuscotra and T. lignorum in uninoculated plants.

In the rhizoplane of JG-62, as a result of spray with indole acetic acid highest increase in the frequency was that of C. echinulata followed by F. oxysporum f.sp. ciceri, A. funigatus, A. flavipes, A. candidus, A. sparsus, B. cinerea, R. solani, A. fusipora, D. hawaiiensis and sterile brown mycelium in inoculated and that of R. oryzae followed by C. aspernum, A. fumigatus, A. flavus, A. niger, A. terreus, A. luchuensis, A. flavipes, T. viridae, A. alternata and sterile black mycelium in uninoculated plants; with indole butyric acid highest increase in the frequency was that of R. oryzae followed by C. echinulata, M. phaseolina, F. oxysporum f.sp. ciceri, A. fumigatus, A. flavus, C. lunata and sterile brown mycelium in inoculated and that of A. fumigatus followed by C. herbarum, M. echinata, R. fuscotra, C. flavum, T. viridae, D. hawaiiensis, C. lunata and sterile black mycelium in uninoculated plants; with thio indole butyric acid R. oryzae followed by A. alternata, S. rolfsii, A. fusipora, A. fumigatus, A. flavus and sterile brown mycelium, M. phaseolina, B. cinerea, C. lunata and F. oxysporum f.sp. ciceri in inoculated and that of A. fumigatus followed by H. fuscotra, C. aspernum, D. hawaiiensis, C. lunata, M. echinata, C. herbarum, T. viridae

and A. flavus in uninoculated plants, with gibberellic acid M. phaseolina followed by M. alpina, R. solani, D. hawaiiensis, A. alternata, C. lunata, F. oxysporum f.sp. ciceri, A. fumigatus, A. flavus and sterile brown mycelium in inoculated and that of A. niger followed by A. alternata, C. lunata, M. echinata, C. herbarum, T. viridae, C. aspermum, A. fumigatus and A. flavus in uninoculated plants. But with maleic hydrazide frequency of A. niger decrease highest followed by M. phaseolina, S. rolfsii, R. solani, M. racemosus, C. flavum, C. echinulata, C. herbarum, M. olivaceum, A. alternata and F. oxysporum f.sp. ciceri in inoculated and that of A. flavipes followed by C. lunata, D. hawaiiensis, A. alternata, A. fumigatus, A. luchuensis, A. sparsus, C. herbarum and sterile black mycelium in uninoculated plants (Table 14).

With an other variety JG-74, as a result of spray with indole acetic acid highest increase in the frequency was that of S. rolfsii followed by R. nodosus, M. globosus, C. echinulata, A. niger, F. oxysporum f.sp. ciceri, C. herbarum and sterile brown mycelium in inoculated and that of A. fumigatus followed by A. niger, A. flavus, C. aspermum, T. viridae, A. humicola, D. hawaiiensis, C. lunata, A. sparsus, A. terreus, A. clavatus, M. echinata and sterile black mycelium in uninoculated plants; with indole butyric acid M. alpina followed by R. nodosus, M. globosus, A. fumigatus, A. niger, F. oxysporum f.sp. ciceri, D. hawaiiensis, C. herbarum, and sterile brown mycelium in

inoculated and that of A. fumigatus followed by A. niger,  
A. sparsus, H. fuscotra, R. oryzae, C. aspermum, M. echinata,  
D. hawaiiensis, G. roseum, T. viridae and sterile black  
mycelium in uninoculated plants; with thio-indole butyric  
acid A. fumigatus followed by R. nodosus, T. viridae,  
M. olivaceum, C. lunata, D. hawaiiensis, M. globosus,  
A. fumigatus, A. niger, C. herbarum, F. oxysporum f.sp. ciceri  
and sterile brown mycelium in inoculated plants and that of  
A. fumigatus followed by A. niger, A. flavus, C. herbarum,  
T. viridae, C. lunata, H. fuscotra, R. oryzae and sterile  
black mycelium in uninoculated plants; with gibberellic acid  
R. nodosus followed by M. globosus, A. fumigatus, C. echinulata,  
A. flavipes, C. herbarum, C. lunata, T. viridae, F. oxysporum  
f.sp. ciceri and sterile brown mycelium in inoculated and that  
of R. oryzae followed by A. flavus, A. fumigatus, A. sparsus,  
A. clavatus, A. terreus, C. aspermum, C. herbarum, M. echinata,  
T. viridae, H. fuscotra, D. hawaiiensis and sterile black  
mycelium in uninoculated plants. On the other hand, with  
maleic hydrazide the frequency of most of the fungi decreased  
and the highest decrease in the frequency was that of A.  
alternata followed by S. rolfsii, R. nodosus, C. echinulata,  
M. globosus, A. flavipes, C. lunata, F. oxysporum f.sp. ciceri  
and sterile brown mycelium in inoculated and that of R. nodosus  
followed by A. terreus, H. fuscotra, R. oryzae, A. fumigatus,  
A. clavatus, C. herbarum, M. echinata, D. hawaiiensis,

C. lunata, T. viridae and sterile black mycelium in uninoculated plants (Table 18). Sprays with growth promoting substances resulted in the increase in the frequency of most of the fungi.

#### Fertilizers:

Results presented in Tables 12 and 16 also show that as a result of spray of plants with fertilizers, the number of fungi in the rhizosphere increased with the increase in the number of sprays and the number of fungi in the rhizosphere was also more in those inoculated plants. In amongst the two varieties tested, the highest number of fungi was found in the rhizosphere of JG-62 both in inoculated and those uninoculated. When sprayed with urea, the number of fungi in the rhizosphere of JG-62 was 21, 28, 30 in inoculated as against 20, 25, 27 in uninoculated plants after respective sprays; with muriate of potash 19, 25, 29 in inoculated as against 18, 24, 26 in uninoculated plants. The number of fungi in the rhizosphere of those sprayed with water as control were 19, 26, 29 in inoculated as against 19, 24, 27 in uninoculated plants. With JG-74, the corresponding values were 23, 27, 29 in inoculated as against 20, 26, 27 in uninoculated plants when sprayed with urea and 22, 26, 28 in inoculated as against 19, 25, 27 in uninoculated plants with potash. The number of fungi in the rhizosphere of plants sprayed with water as

control was 21, 26, 28 in inoculated as against 19, 25, 27 in uninoculated plants.

Results with rhizoplane fungi also exhibited a similar trend (Table 14 and 18). In amongst the two cultivars tested number of fungi was more in rhizoplane of variety JG-62 both in inoculated and uninoculated plants. In rhizoplane of inoculated plants of JG-62, when sprayed with urea, the number of fungi was 17, 21, 24 in inoculated as against 15, 19, 20 in respective sprays in uninoculated plants and with muriate of potash 17, 21, 23 in inoculated as against 13, 19, 20 in uninoculated plants. The number of fungi was 15, 20, 23 in inoculated as against 15, 19, 20 in uninoculated plants when sprayed with water. With cv. JG-74 the corresponding values were 15, 18, 20 in inoculated as against 14, 18, 19 in uninoculated plants when sprayed with urea and 15, 17, 20 in inoculated as against 14, 18, 18 in uninoculated plants with Muriate of potash. The number of fungi was 15, 18, 20 in inoculated as against 14, 18, 19 in uninoculated plants when sprayed with water as control. However, the number of fungi was less in rhizoplane as compared to rhizosphere.

The population of fungi in the rhizosphere of both the varieties increased due to spray with urea however with muriate of potash, there was a decrease in the rhizosphere of JG-62 (Tables 10 and 11). However, the increase was more in inoculated

ones as compared to uninoculated ones.

Frequency of fungi in general showed an increase as a result of spray with urea. The frequency of sterile brown mycelium was highest in the rhizosphere of inoculated plants of JG-62 followed by Rhizopus oryzae, Monosporium olivacium, Botrytis cinerea, Spicaria silvatica and Fusarium oxysporum f.sp. ciceri that of R. oryzae, followed by Chaetomium flavum, S. silvatica, Memnoniella echinata, Fusarium moniliformae and sterile black mycelium in uninoculated; with muriate of potash, R. oryzae followed by Aspergillus fumigatus, A. flavus, A. niger, in inoculated plants as against Trichoderma viridae followed by sterile black mycelium, M. echinata, Aspergillus terreus, A. sparsus and R. oryzae in uninoculated plants (Table 12). In the rhizosphere of another variety JG-74, the highest increase in the frequency was that of T. viridae followed by sterile brown mycelium, A. flavus, A. niger, A. flavipes, M. echinata, Rhizopus nodosus, Phoma hibernica, Gliocladium roseum, Penicillium notatum, Alternaria alternaria and F. oxysporum f.sp. ciceri when sprayed with urea in inoculated and that of Cephalosporium aspermum followed by A. fumigatus. A. niger, A. flavus, A. terreus, A. clavatus, F. moniliformae, Rhizopus nodosus, T. viridae, M. echinata, Cladosporium herbarum, Alternaria humicola, G. roseum and P. notatum in uninoculated plants; with muriate of potash highest increase in the frequency was that of sterile



brown mycelium followed by M. echinata, T. viridae, A. clavatus, A. niger, A. flavus, A. fumigatus and R. nodosus in inoculated and that of sterile black mycelium followed by R. nodosus, T. viridae, A. clavatus, A. niger, A. fumigatus, A. terreus, P. hibernica and M. echinata in uninoculated plants (Table 16).

A similar trend was observed with rhizoplane fungi (Tables 14 and 18). As a result of spray with urea, highest increase in the frequency was that of Cunninghamella echinulata, followed by A. flavus, A. fumigatus, A. niger, sterile brown mycelium, C. flavum, D. hawaiiensis, R. oryzae, S. rolfsii and F. oxysporum f.sp. ciceri in the rhizoplane of inoculated plants of JG-62; and A. fumigatus followed by A. flavus, A. niger, A. terreus, Hemicola fuscotra, Curvularia lunata, C. herbarum, M. echinata, T. viridae, R. oryzae, C. asperum and sterile black mycelium in uninoculated plants, with muriate of potash highest increase in the frequency was that of A. flavus followed by A. niger, A. fumigatus, R. oryzae, C. echinulata, M. racemosus, M. alpina, C. herbarum and sterile brown mycelium in inoculated and A. terreus followed by A. flavus, A. niger, A. fumigatus, M. echinata, H. fuscotra, T. viridae and sterile black mycelium in uninoculated plants. With another variety JG-74 as a result of spray with urea highest increase in the frequency was that of A. clavatus followed by R. nodosus, T. viridae, sterile brown mycelium, A. flavus, A. fumigatus, C. echinulata,

M. alpina, D. hawaiiensis, M. olivaceum, S. rolfsii and F. oxysporum f.sp. ciceri in inoculated and A. fumigatus followed by A. flavus, A. niger, A. terreus, A. clavatus, R. oryzae, C. aspermum, G. roseum, M. echinata, T. viridae, H. fuscotra and A. humicola in uninoculated plants, with muriate of potash highest increase in the frequency was that of A. clavatus followed by T. viridae, R. nodosus, A. fumigatus, A. flavus, A. niger, G. roseum and sterile brown mycelium in inoculated and R. nodosus followed by A. sparsus, A. terreus, A. clavatus, A. funigatus, T. viridae, M. echinata, D. hawaiiensis and sterile black mycelium in uninoculated plants.

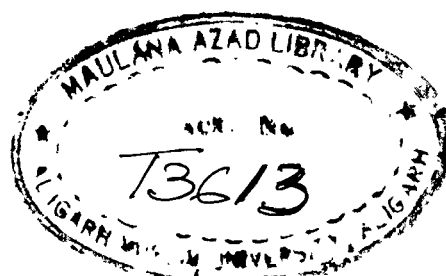
#### Pesticides:

Results presented in Tables 13 and 17 reveal that due to spray of plants with pesticides, the number of fungi in the rhizosphere increased with increase in the number of sprays but this increase was low<sup>er</sup> than those sprayed with water. The number of fungi was also more in those inoculated with Fusarium oxysporum f.sp. ciceri. In amongst the two varieties tested the more number of fungi was reported from the rhizosphere of JG-62 both in inoculated and uninoculated when sprayed with bavistin the number of fungi in cv. JG-62 was 19, 24, 27 in inoculated as against 18, 24, 25 in uninoculated plants after respective sprays; with brassicol 15, 23, 27 in inoculated as against 15, 21, 23 in uninoculated plants; with vitavax 16, 24, 27

in inoculated as against 17, 22, 25 in uninoculated plants; with benlate, 14, 22, 24 in inoculated as against 14, 21, 23 in uninoculated plants; with captan 15, 23, 27 in inoculated as against 17, 22, 24 in uninoculated plants, with wettable sulphur 18, 24, 25 in inoculated as against 18, 23, 25 in uninoculated plants; and with streptomycine 20, 26, 29 in inoculated as against 19, 25, 27 in uninoculated plants. The number of fungi was 19, 26, 29 inoculated and 19, 24, 27 in uninoculated plants. when sprayed with water. With JG-74 the corresponding values were 22, 25, 26 in inoculated as against 19, 23, 24 in uninoculated plants when sprayed with bavistin, 22, 25, 25 as against 17, 21, 20 with brassicol, 22, 25, 24 as against 17, 23, 24 with vitatax, 20, 24, 24 as against 18, 21, 21 with benlate, 21, 24, 25 as against 18, 22, 23 with captan, 22, 25, 25 as against 17, 21, 22 with wettable sulphur and 22, 26, 26 as against 20, 23, 24 with streptomycine. The number of fungi in the rhizosphere of plants sprayed with water as control was 21, 25, 28 in inoculated as against 19, 25, 27 in uninoculated plants. An increase in the number of fungi in the rhizosphere was due to streptomycine sprays which was more than those spryaed with water out in the remaining there was decrease over control.

Almost identical results were observed with reaged to rhizoplane (Tables 15 and 19). Here also there <sup>have</sup> been more fungi in those inoculated with F. oxysporum f.sp. ciceri and

that too in JG-62. When sprayed with bavistin the number of fungi in rhizoplane of inoculated JG-62 plants was 16, 20, 21 as against 13, 19, 20 in respective sprays in uninoculated; with brassicol 15, 20, 23 in inoculated as against 12, 18, 19 in uninoculated; with vitavax 14, 20, 20 in inoculated as against 13, 19, 20 in uninoculated; with benlate 15, 20, 21 in inoculated as against 12, 18, 18 in uninoculated plants; with captan 13, 20, 21 in inoculated as against 12, 19, 21 in uninoculated plants; with wettable sulphur 17, 20, 22 in inoculated as against 13, 14, 19 in uninoculated plants; with streptomycin 16, 21, 23 in inoculated as against 13, 19, 19 in uninoculated plants. While the number of fungi in plants sprayed with water as control was 15, 20, 23 in inoculated as against 15, 19, 20 in uninoculated after respective sprays. With JG-74 the corresponding values were 14, 16, 18 in inoculated as against 14, 18, 18 in uninoculated when sprayed with bavistin; 15, 18, 19 as against 14, 18, 17 with brassicol 15, 18, 18 as against 14, 18, 18 with vitavax, 15, 15, 17 as against 14, 18, 18 with benlate, 14, 17, 18 as against 14, 18, 18 with captan, 14, 17, 18 as against 13, 17, 17 with wettable sulphur and 17, 19, 20 as against 16, 19, 19 with streptomycin. The number of fungi in the rhizoplane of plants sprayed with water as control was 15, 18, 20 in inoculated as against 14, 18, 19 in uninoculated plants. Here also the highest increase in the number of fungi was due to spray with streptomycine. However the number of fungi was less in rhizoplane as



compared to rhizosphere.

By and large the population of fungi in the rhizosphere of inoculated and uninoculated plants of both the varieties decreased due to spray with pesticides (Tables 10 and 11). The spray with streptomycin however brought about an increase. Highest decrease was observed when sprayed with bavistin and benlate.

Frequency of fungi in general showed a decrease as a result of spray with pesticides (Tables 13 and 17). As a result of spray with bavistin the highest reduction in the frequency in the rhizosphere of JG-62 inoculated plants was that of Fusarium oxysporum f.sp. ciceri followed by Mortierella alpina, Aspergillus flavipes, Curvularia lunata, Alternaria alternata, Monosporium olivaceum, Monilia humicola that of Cephalosporium asperum followed by Aspergillus sparsus, Torulla allii, Memnoniella echinata, C. lunata, C. geniculata, Acrophialophora fusipora, Drechslera hawaiiensis, Cladosporium herbarum, Alternaria alternata and Aspergillus terreus in uninoculated; with vitavax A. sparsus followed by Botrytis cinerea, Sclerotium rolfsii, Rhizoctonia solani, Syncephalastrum racemosum, Mucor racemosus, Aspergillus candidus, A. alternata and F. oxysporum f.sp. ciceri, B. silvatica, A. alternata, C. flavum and sterile brown mycelium in inoculated and that of A. terreus followed by T. allii, H. fuscotra,

C. geniculata, A. alternata, T. viridae, C. herbarum, A. niger,  
A. luchuensis, F. moniliiformae, C. flavum, R. oryzae,  
M. humicola and sterile black mycelium in uninoculated plants,  
with brassicol, B. cineria followed by C. flavum, R. solani,  
C. lunata, S. silvatica, A. flavipes, A. sparsus, A. candidus,  
S. chrysospermum and A. fusipora in inoculated and that of  
A. terreus followed by A. flavipes, A. funiculosus, A. luchuensis,  
A. sparsus, A. alternata, F. moniliiformae, C. aspermum,  
R. oryzae and sterile black mycelium in uninoculated plants,  
with benlate F. oxysporum f.sp. ciceri followed by sterile  
brown mycelium, M. alpina, A. alternata, M. racemosus,  
M. phaseolina, A. candidus, A. flavipes, A. sparsus, S. rolfsii,  
R. solani and B. cineria in inoculated and that of T. allii  
followed by C. aspermum, A. terreus, A. flavipes, H. fuscotra,  
C. lunata, C. geniculata, A. fusipora, M. humicola, C. flavum,  
F. moniliiformae and sterile black mycelium in uninoculated  
plants; with captan A. candidus followed by C. flavum,  
M. phaseolina, S. rolfsii, F. oxysporum f.sp. ciceri,  
S. chrysospermum, A. alternata, C. lunata, A. fusipora,  
A. flavipes, S. racemosus, M. olivaceum and B. cineria  
in inoculated and that of T. allii followed by B. pulluliferum,  
A. terreus, A. funiculosus, A. sparsus, A. flavipes,  
F. moniliiformae, S. silvatica, T. viridae and sterile black  
mycelium in uninoculated plants; with wettable sulphur  
S. rolfsii followed by C. lunata, A. alternata, C. flavum,

M. olivaceum, B. cinerea, S. racemosum, M. phaseolina,  
A. sparsus, A. flavipes, S. chrysospermum, R. solani and  
F. oxysporum f.sp. ciceri in inoculated and that of A. sparsus  
followed by A. flavipes, H. fuscotra followed by C. lunata,  
F. moniliiformae, T. viridae, D. hawaiiensis, T. allii,  
A. terreus and A. alternata in uninoculated plants. On the  
other hand, with streptomycine the frequency of most of the  
fungi increased with highest increase that of sterile brown  
mycelium followed by M. racemosus, R. oryzae, M. humicola,  
A. fumigatus, A. sparsus, B. cinerea, R. solani and F. oxysporum  
f.sp. ciceri in inoculated and that of sterile black mycelium  
followed by B. piluliferum, C. aspermum, M. echinata,  
C. lunata, F. moniliiformae, T. viridae, C. herbarum, M. humicola  
and R. oryzae in uninoculated plants. With another variety  
JG-74, as a result of spray with bavistin the highest reduction  
in the frequency was that of F. oxysporum f.sp. ciceri followed  
by S. rolfsii, A. flavipes, M. olivaceum, A. alternata,  
C. lunata, P. notatum, M. alpina and sterile brown mycelium  
in the rhizosphere of inoculated plants and that of R. nodosus  
followed by C. aspermum, P. hibernica, T. allii, C. lunata,  
F. moniliiformae, H. fuscotra, D. hawaiiensis, D. sativum,  
C. lunata, A. humicola, T. lignorum, T. viridae and sterile  
black mycelium in uninoculated plants, with vitavax, V. glaucum,  
followed by A. alternata, S. rolfsii, C. lunata, A. candidus,  
M. alpina, C. flavum, M. humicola, P. notatum, G. roseum

and sterile brown mycelium in inoculated plants and that of D. sativum followed by T. lignorum, T. viridae, A. humicola, C. lunata, H. fuscotra, G. herbarum, T. allii, R. nodosus, C. aspernum, P. hibernica, A. clavatus and sterile brown mycelium in uninoculated plants; with brassicol that of C. lunata followed by V. glaucum, S. rolfsii, T. viridae, T. lignorum, P. notatum, M. olivaceum, G. racemosum and F. oxysporum f.sp. ciceri in inoculated and that of A. clavatus followed by A. humicola, C. lunata, R. nodosus, T. lignorum and M. echinata in uninoculated plants; with benlate that of F. oxysporum f.sp. ciceri followed by V. glaucum, A. candidus, A. clavatus, A. alternata, T. viridae, D. hawaiiensis, P. notatum, G. racemosum, M. olivaceum, C. lunata and sterile brown mycelium in inoculated and that of R. nodosus, followed by P. hibernica, A. clavatus, P. notatum, G. roseum, T. allii, C. herbarum, M. echinata, D. hawaiiensis, A. humicola, F. moniliiformae, T. lignorum and sterile black mycelium in uninoculated plants; with captan that of F. oxysporum f.sp. ciceri followed by T. lignorum, V. glaucum, A. candidus, A. clavatus, M. alpina and A. alternata inoculated and that of F. moniliiformae followed by A. humicola, C. lunata, D. hawaiiensis, T. allii, P. notatum, A. clavatus, A. sparsus, C. aspernum and sterile black mycelium in uninoculated plants; with wettable sulphur C. echinulata followed by M. globosus, M. alpina, V. glaucum, C. lunata and F. oxysporum f.sp. ciceri in inoculated and that of R. nodosus



followed by P. hibernica, A. terreus, A. clavatus, P. notatum, M. echinata, D. hawaiiensis, D. sativum, C. lunata, A. humicola, T. lignorum and sterile black mycelium in uninoculated plants. On the other hand, with streptomycine the frequency of fungi increased with highest increase that of A. clavatus followed by A. flavus, A. fumigatus, A. niger, A. flavipes, A. candidus, A. alternata, P. notatum, C. flavum, R. nodosus, M. echinata, T. viridae and sterile brown mycelium in inoculated and that of P. hibernica followed by A. fumigatus, A. flavus, A. niger, A. sparsus, A. flavipes, C. herbarum, M. echinata, H. fuscotra, T. viridae, T. lignorum, C. asperum, P. notatum, G. racemosum and sterile black mycelium in uninoculated plants.

A similar trend was observed the changes in frequency of rhizoplane fungi (Tables 15 and 19). As a result of spray with bavistin highest decrease in the frequency was that of F. oxysporum f.sp.ciceri followed by S. rolfsii, A. alternata, B. cinerea, A. flavipes, M. phaseolina, C. lunata, D. hawaiiensis, A. sparsus, A. candidus, M. albina and sterile brown mycelium in inoculated plants and that of A. fusipora followed by M. echinata, C. lunata, A. terreus, A. flavus, A. sparsus, A. flavipes, H. fuscotra, T. viridae and sterile black mycelium in uninoculated plants; with vitavax B. cinerea followed by M. albina, A. alternata, C. lunata, M. phaseolina, C. flavum, A. flavus, A. sparsus and sterile brown mycelium in inoculated and that of C. asperum followed by A. terreus, A. luchuensis,

A. humicola, D. hawaiiensis, A. flavus, A. sparsus, A. flavipes,  
A. niger, R. oryzae, H. fuscotra, T. viridae, C. herbarum,  
M. echinata and sterile black mycelium in uninoculated plants;  
with brassicol A. flavipes followed by S. rolfsii, A. fusipora,  
A. alternata, C. lunata, R. solani, A. sparsus, A. candidus,  
B. cinerea and sterile brown mycelium in inoculated and that  
of H. fuscotra followed by C. lunata, C. herbarum, M. echinata  
A. terreus, A. niger, A. flavipes, A. sparsus, A. luchuensis,  
D. hawaiiensis, T. viridae and sterile black mycelium in  
uninoculated plants, with benlate F. oxysporum f.sp. ciceri  
followed by M. olivaceum, A. fusipora, C. lunata, R. oryzae,  
A. candidus, A. flavipes, M. racemosus, M. alpina, C. flavum  
M. phaseolina, B. cinerea, A. alternata and sterile brown  
mycelium in inoculated and that of M. echinata followed by  
T. viridae, A. niger, A. terreus, A. flavus, D. hawaiiensis,  
C. lunata, C. aspermum, R. oryzae, C. herbarum, C. flavum and  
sterile black mycelium in uninoculated plants; with captan  
B. cinerea followed by C. flavum, M. phaseolina, A. alternata,  
M. alpina, C. lunata, M. olivaceum, M. phaseolina, R. solani,  
S. rolfsii, A. candidus, A. flavipes, A. sparsus, F. oxysporum  
f.sp. ciceri and sterile brown mycelium in inoculated and that  
of A. flavipes followed by A. terreus, A. sparsus, C. herbarum,  
D. hawaiiensis, C. lunata, S. rolfsii, R. oryzae, C. aspermum,  
A. niger, T. viridae, H. fuscotra and sterile black mycelium  
in uninoculated plants, on the other hand with streptomycin

the frequency of most of the fungi increased in both inoculated and uninoculated plants. The frequency of M. phaseolina increased highest followed by A. candidus, A. flavipes, A. sparsus, A. fumigatus, A. flavus, R. oryzae, M. racemosus, M. alpina, C. lunata, D. hawaiiensis, S. rolfsii, R. solani, B. cinerea, F. oxysporum f.sp. ciceri in inoculated and that of A. fumigatus followed by C. lunata, C. herbarum, T. viridae, D. hawaiiensis, R. oryzae, C. aspermum, A. niger, A. flavipes, M. echinata, Fusarium oxysporum and sterile black mycelium in uninoculated plants. With another variety JG-74, as a result of spray with bavistin highest reduction in the frequency was that of A. alternata in the rhizoplane of inoculated plants followed by S. rolfsii, M. olivaceum, G. roseum, F. oxysporum f.sp. ciceri, M. alpina and sterile brown mycelium and that of R. oryzae followed by A. terreus, M. echinata, T. viridae, C. aspermum, A. flavus, A. clavatus, A. sparsus, H. fuscotra, D. hawaiiensis, A. humicola, C. lunata, C. herbarum and sterile black mycelium in uninoculated plants; with vitavax A. alternata followed by F. oxysporum f.sp. ciceri, M. alpina, G. roseum, S. rolfsii, C. lunata and sterile brown mycelium in inoculated and that of A. humicola followed by C. herbarum, R. nodosus, D. hawaiiensis, C. lunata, H. fuscotra, T. viridae, C. aspermum, A. flavus, A. terreus, A. sparsus and sterile black mycelium in uninoculated plants; with brassicol M. alpina followed by A. alternata, F. oxysporum f.sp. ciceri, A. flavipes,

G. roseum , T. viridae and sterile brown mycelium in inoculated and that of R. nodosus followed by D. hawaiiensis, T. viridae M. echinata, C. aspermum, A. terreus, A. flavus, A. clavatus, A. sparsus, R. oryzae, C. lunata and A. humicola in uninoculated plants; with benlate F. oxysporum f.sp. ciceri followed by C. lunata, M. olivaceum, S. rolfsii, A. clavatus, A. flavipes, C. echinulata, M. globosus, G. roseum , T. viridae and sterile brown mycelium in inoculated and that of A. clavatus followed by H. fuscotra, M. echinata, R. nodosus, C. herbarum, G. roseum , T. viridae, A. terreus, A. flavus, R. oryzae, A. humicola, D. hawaiiensis, C. lunata and sterile black mycelium in uninoculated plants; with captan M. alpina followed by F. oxysporum f.sp. ciceri, C. lunata, C. echinulata, A. alternata, A. clavatus in inoculated and that of R. nodosus followed by A. humicola, A. clavatus, C. herbarum, A. niger, A. terreus A. sparsus, R. oryzae, H. fuscotra, T. viridae, M. echinata, C. aspermum, and D. hawaiiensis in uninoculated plants; with wettable sulphur R. nodosus followed by M. alpina, M. olivaceum, F. oxysporum f.sp. ciceri, C. echinulata, M. globosus, A. flavipes, A. alternata and C. lunata in inoculated and that of A. clavatus followed by R. nodosus, A. terreus, A. sparsus, A. humicola, C. lunata, M. echinata, R. oryzae H. fuscotra, A. niger, A. flavus, D. hawaiiensis, G. roseum and sterile black mycelium in uninoculated plants. On the other hand with streptomycin the frequency of fungi increased

with highest increase that of R. nodosus followed by A. clavatus, A. alternata, C. echinulata, A. flavipes, M. olivaceum, T. viridae, M. echinata and sterile brown mycelium in inoculated and that of A. fumigatus followed by A. niger, A. flavus, A. clavatus, A. terreus, A. humicola, R. nodosus, R. oryzae, C. aspermum, G. roseum , M. echinata, T. viridae, D. hawaiiensis, C. lunat and sterile black mycelium in uninoculated plants.

It is interesting that as a result of spray with growth promoting substances, there has been higher increase in the frequency of saprophytic fungi in the rhizosphere of both the varieties except with gibberellic acid where highest increase was that of Fusarium oxysporum f.sp. ciceri a pathogenic fungi in inoculated JG-62 plants. It might be one of the factor for high incidence of wilt disease due to spray with gibberellic acid. This was not true however with the rhizosphere of JG-74 which is relatively less susceptible to wilt. There were higher number of saprophytic fungi in the treatment with indole butyric acid followed by gibberellic acid, thio indole butyric acid and indole acetic acid with the rhizosphere of JG-62; but with the rhizosphere of JG-74 it was gibberellic acid, thio indole butyric acid indole acetic acid and indole butyric acid. On the other hand with maleic hydrazide, there was a reduction in the frequency of large number of saprophytic, parasitic and also some antibiotic producing fungi in both the varieties.

With rhizoplane fungi the frequency of fungi showing increase particularly the saprophytic fungi in the rhizosphere of JG-62 was more with indole butyric acid followed by gibberellic acid, indole acetic acid and thio indole butyric acid. It appears that indole butyric acid was more favourable for increase in saprophytic fungi however with JG-74 gibberellic acid was more effective followed by thio indole butyric acid. The response of indole butyric acid and indole acetic acid was almost identical. On the other hand with maleic hydrazide, generally the frequency of fungi showed reduction and this reduction was more in the saprophytic fungi in both the cultivars.

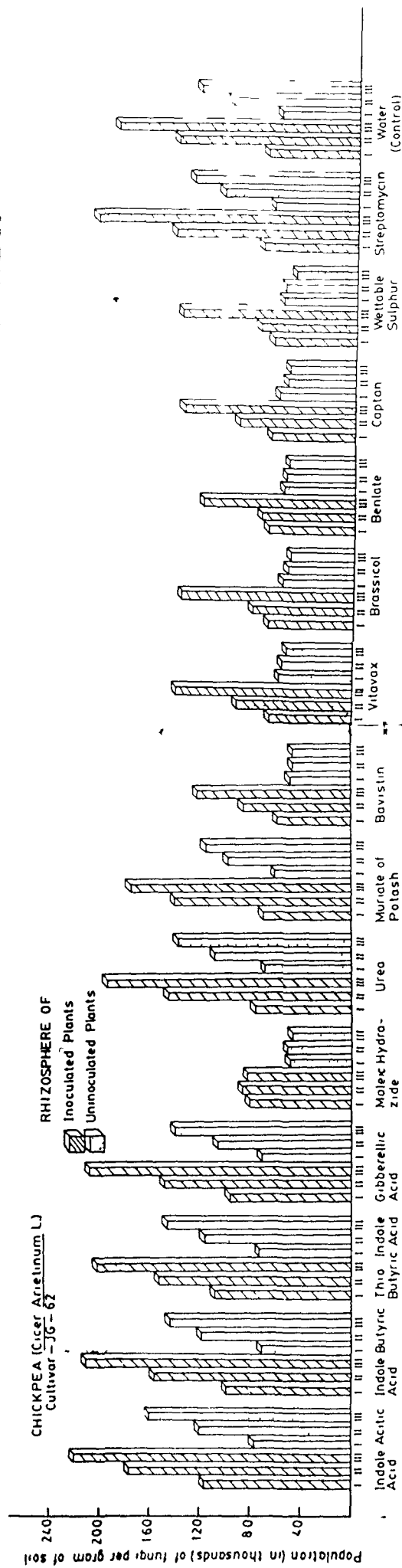
With fertilizers urea was more effective than muriate of potash in increasing the activity of saprophytic fungi but with muriate of potash highest increase in the frequency was that of T. viridae in the rhizosphere of JG-74 inoculated plants which could be exploited for control. Similarly with cv. JG-74 T. viridae was greatly increased with muriate of potash and urea in inoculated and urea followed by muriate of potash in uninoculated plants.

In the rhizoplane also urea increased the frequency of large number of saprophytic fungi in both the varieties in inoculated and muriate of potash in uninoculated of cv. JG-62 but in cv. JG-74 effect of both the fertilizers was identical.

As a result of spray with fungicides there was a reduction in the frequency of large number of saprophytic and parasitic fungi. In some cases even the frequency of antibiotic producing fungi decreased. The spray with streptomycin however brought about an increase in frequency of all the fungi.

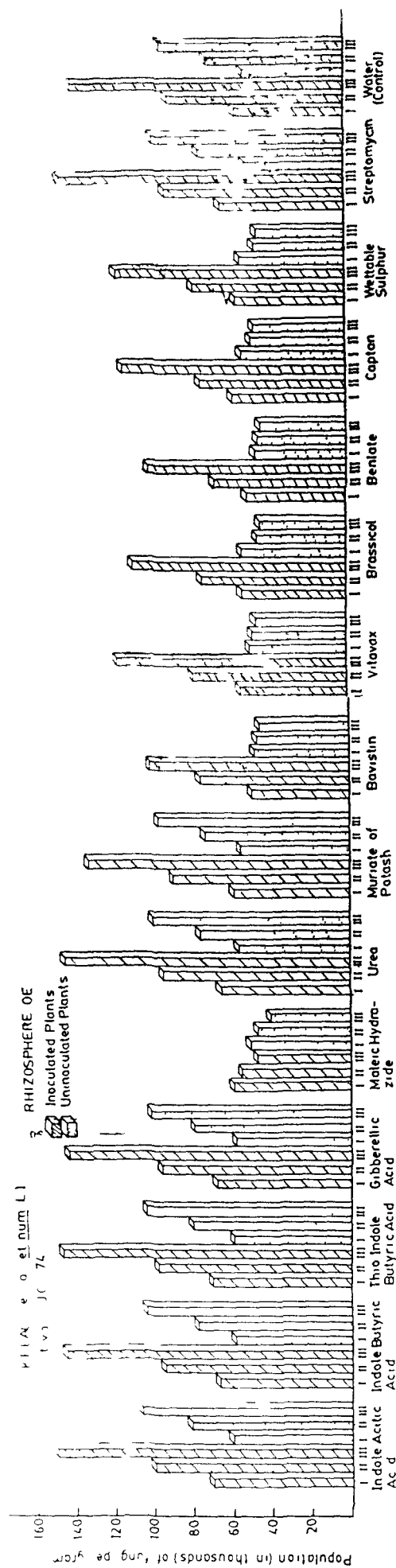
FIG. 3. Population of fungi in the rhizosphere of uninoculated and inoculated plants of chickpea cultivar JG-62 (with Fusarium oxysporum f.sp. ciceri) treated with different foliar sprays.





FOLIAR SPRAY  
Fig 3

FIG. 4. Population of fungi in the rhizosphere of uninoculated and inoculated plants of chickpea cultivar JG-74 (with Fusarium oxysporum f.sp. ciceri) treated with different foliar sprays.



FOLIAR SPRAY  
Fig 4

TABLE - 10: Effect of foliar spray of different chemicals on population of fungi (per gram of soil) in the non-rhizosphere and rhizosphere cultivar JG-62 uninoculated and inoculated with Fusarium oxysporum f.sp. ciceri

Plant Material	JG-62					
	INOCULATED PLANTS			UNINOCULATED PLANTS		
	I Spray	R:S	II Spray	R:S	III Spray	R:S
Non-rhizosphere	15.5		16.5		17.0	
Control	72.5	4.68	142.5	8.64	190.0	11.17
Indole Acetic Acid	115.0	7.42	176.0	10.67	220.0	12.94
Indole Butyric Acid	98.0	6.32	155.0	9.39	210.0	12.35
Thio-indole Butyric Acid	108.0	6.67	152.0	9.21	202.0	11.88
Gibberellic Acid	95.0	6.13	148.0	8.97	208.0	12.24
Maleic Hydrazide	80.0	5.16	85.5	5.18	82.5	4.85
Urea	76.0	4.90	145.0	8.79	193.0	11.35
Muriate of Potash	70.0	4.52	140.0	6.06	125.0	7.39
Baustein	58.0	3.74	85.0	5.15	122.0	7.18
Vitavax	66.5	4.29	92.0	4.36	140.0	8.24
Brassiccol	68.5	4.42	80.5	4.27	137.0	8.06
Benlate	68.0	4.39	74.0	4.48	120.0	7.06
Captan	67.0	4.32	93.5	5.67	135.0	7.94
Wettable Sulphur	65.0	4.19	75.0	4.55	138.0	8.12
Streptomycine	74.0	4.77	144.5	8.72	205.5	12.05
L.S.D. at 5%	1.4467968		1.5512628		1.647438	
L.S.D. at 1%	1.9484286		2.089115		2.2186959	
				0.8593665	0.8496069	0.8517855
				1.1573251	1.1441817	1.1471157

In thousands?

Needs analysis

TABLE - 11: Effect of foliar spray of different chemicals on population of *Fus. i* (per gram of soil) in the non-rhizosphere and rhizosphere of chickpea cv JG-74 uninoculated and inoculated with *Fusarium oxysporum* f.sp. *ciceri*

TREATMENT	JG-74									
	INOCULATED PLANTS					UNINOCULATED PLANTS				
	I Spray	R:S	II Spray	R:S	III Spray	R:S	I Spray	R:S	II Spray	III Spray
Non-rhizosphere	15.5		16.5		17.0		15.5		16.5	17.0
Control	56.0	3.61	90.0	5.45	140.0	8.24	52.0	3.35	70.0	95.0
Indole Acetic Acid	71.0	4.73	100.0	6.06	150.0	8.82	60.0	3.87	81.0	107.0
Indole Butyric Acid	66.0	4.26	95.0	5.76	145.0	8.53	58.5	3.77	77.5	103.5
Thio-indole Butyric Acid	70.0	4.52	98.0	5.94	147.0	8.64	59.0	3.81	80.0	104.0
Gibberellic Acid	68.5	4.41	96.0	5.82	144.0	8.47	58.0	3.74	79.0	102.0
Maleic Hydrazide	59.0	3.81	55.0	3.34	51.5	3.02	50.0	3.23	46.0	40.0
Urea	65.0	4.20	94.5	5.73	145.0	8.53	56.0	3.61	76.0	100.0
Muriate of Potash	58.0	3.74	90.0	4.85	134.5	7.91	55.0	3.55	74.0	98.0
Bavistin	49.0	3.16	76.0	4.61	102.0	6.00	47.0	3.03	46.5	45.0
Vitavax	55.0	3.55	78.5	4.76	118.0	6.94	49.5	3.19	47.5	46.5
Brassiccol	54.0	3.48	75.5	4.56	110.0	6.47	53.5	3.42	46.0	44.0
Benlate	51.5	3.32	68.0	4.12	102.0	6.00	46.0	2.97	45.5	43.5
Captan	58.0	3.74	76.0	4.61	115.0	6.76	53.0	3.42	48.5	47.0
Wettable Sulphur	56.5	3.66	78.0	4.73	117.0	6.88	54.0	3.48	46.0	45.0
Streptomycine	64.0	4.13	92.0	5.58	146.0	8.58	55.5	3.58	74.5	99.0
L.S.D. at 5%	1.8375476		2.0939766		1.934818		1.4560538		1.5805541	2.5850337
L.S.D. at 1%	2.4746601		2.8199979		2.605656		1.9608952		2.1285621	3.5157482

needs analysis











TABLE - 15: Frequency (percentage) of fungi in the rhizosphere of chickpea plants cultivated with Fusarium oxysporum f.sp. olicei sprayed with pesticide.

Fungi isolated	CONTROL						BAYISTIN						VITAVAX						BRASSICOL						BENLATE						CAPTAN						WETTABLE SULPHUR						ZEPHYRUS																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																												
	IP			UP			IP			UP			IP			UP			IP			UP			IP			UP			IP			UP			IP			UP			IP			UP																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																									
	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I

IP = Inoculated plants, CP = uninoculated plants.

+Number of 11,

TABLE - 17: Frequency of fungi in the rhizosphere of chickpea plants cultivar JG-74 uninoculated and inoculated with Fusarium oxysporum f.sp. ciceri sprayed with pesticides

Fungal isolated	CONTROL						BAYVISTIN						VITAVAX						BRASSICOL						BENLATE						CAPTAN						ASTABLE SULPHUR						STREPTOMYCIN																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																				
	IP			UP			IP			UP			IP			UP			IP			UP			IP			UP			IP			UP			IP			UP			IP			UP																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																																	
	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III

+Number of spray

IP = Inoculated plants, UP = Uninoculated plants

TABLE - 13: Frequency (percentage) of fungi in the rhizoplane of chickpea plants cultivated with Fusarium oxysporum f.sp. ciceri sprayed with growth regulators and fertilizers

Fungi isolated	CONTROL			INDOLE ACETIC ACID			INDOLE BUTYRIC ACID			THIO INDOLE BUTYRIC ACID			GIBBERELIC ACID			MALEIC HYDRAZIDE			UREA			MURIATE OF POTASH		
	IP			UP			IP			IP			IP			IP			IP			IP		
	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III	I	II	III
<i>Rhizopus oryzae</i>	-	-	-	20	25	30	-	-	-	-	-	-	25	35	45	-	-	-	20	20	15	-	-	-
<i>R. nodosus</i>	15	10	10	15	25	30	20	25	20	25	20	25	15	25	35	10	5	-	15	10	5	25	20	25
<i>Cunninghamella echinulata</i>	15	20	20	-	-	-	25	25	25	25	20	25	-	-	-	10	15	10	-	-	-	20	25	30
<i>Mucor globosus</i>	10	15	20	-	-	-	15	20	30	-	-	-	20	30	40	-	-	-	-	-	-	15	25	30
<i>Mortierella alpina</i>	10	15	20	-	-	-	20	25	35	-	-	-	20	25	30	-	-	-	-	-	-	10	10	15
<i>Cephalosporium asperum</i>	-	-	-	25	30	50	-	-	-	-	-	-	-	-	-	-	-	-	40	50	60	-	-	-
<i>Aspergillus fumigatus</i>	50	35	30	40	30	40	70	40	35	65	50	45	55	55	55	50	40	35	35	25	10	60	55	60
<i>A. flavus</i>	40	35	35	35	40	45	60	60	60	45	50	50	40	40	70	35	30	25	45	50	55	50	50	55
<i>A. niger</i>	20	25	40	30	35	50	30	40	50	35	40	60	45	60	60	25	35	40	30	40	55	25	35	45
<i>A. terreus</i>	-	-	-	25	35	45	-	-	-	-	-	-	35	45	50	-	-	-	30	20	10	-	-	-
<i>A. clavatus</i>	20	15	15	20	30	40	20	25	25	25	20	20	25	35	45	15	10	10	15	20	25	25	30	35
<i>A. sparsus</i>	-	-	-	15	20	20	-	-	-	-	-	-	-	-	30	-	-	-	10	25	35	-	-	-
<i>A. flavipes</i>	25	20	20	-	-	-	25	20	10	40	35	25	-	-	-	15	10	10	-	-	-	25	30	30
<i>Glodadium roseum</i>	20	25	30	25	30	40	25	30	35	20	25	40	25	30	25	10	15	20	-	-	15	20	25	35
<i>Monosporium olivaceum</i>	-	10	15	-	-	-	-	10	20	-	15	20	-	-	-	-	-	10	-	-	-	-	-	-
<i>Glaesporium herbarum</i>	30	35	45	20	25	35	35	40	50	30	40	50	25	35	45	25	20	25	25	20	15	30	40	30
<i>Nemnoniella echinata</i>	-	-	20	25	30	40	-	-	25	-	-	25	30	35	45	-	-	20	25	20	20	-	15	35
<i>Trichoderma viridae</i>	25	20	15	25	30	40	25	30	25	25	25	25	30	35	50	25	10	10	30	25	25	35	30	35
<i>Humicola fuscotra</i>	-	-	-	-	15	20	-	-	-	-	-	-	-	-	30	-	-	-	-	10	5	-	-	-
<i>Fusarium oxysporum f.sp. ciceri</i>	30	35	40	-	-	-	30	40	50	35	40	45	-	-	-	25	30	10	-	-	-	35	40	50
<i>Alternaria alternata</i>	10	15	20	-	-	-	10	20	25	10	20	25	15	25	35	-	5	5	-	-	-	10	20	25
<i>A. humicola</i>	-	-	-	-	10	20	-	-	-	-	-	-	-	-	-	-	-	-	-	15	15	-	-	-
<i>Drechslera hawaiiensis</i>	-	30	20	-	15	20	-	15	30	-	35	30	-	20	25	-	25	20	-	30	30	-	30	25
<i>Curvularia lunata</i>	-	30	40	-	20	30	-	30	40	-	35	60	-	25	40	-	15	20	-	25	35	-	25	30
<i>Sclerotium rolfsii</i>	-	-	15	-	-	15	-	15	20	-	10	20	-	-	20	-	-	5	-	-	-	-	-	5
Sterile black mycelium	-	-	-	25	30	35	-	-	-	-	-	-	30	35	50	-	-	-	25	25	20	-	-	-
Sterile brown mycelium	25	30	35	-	-	-	35	40	45	-	40	45	50	-	-	40	20	-	-	-	25	30	35	40

\*Number of spray  
IP = Inoculated plants, UP = Uninoculated plants

TABLE 19: Frequency (percentage) of fungi in the rhizosphere of chickpea plants cultivated with *Fusarium oxysporum* f.sp. *ciceri* sprayed with pesticides

[illegible]

† 11 mb 1 of 3ray

3.4 RHIZOSPHERE AND RHIZOPLANE MYCOFLORA OF CHICKPEA PLANTS  
CULTIVAR JG-62 AND JG-74 INOCULATED WITH FUSARIUM  
OXYSPORUM F.SP. CICERI IN RELATION TO SOIL AMENDMENT

Result presented in Table 21 and 23 reveal that soil amendment influenced the rhizosphere mycoflora both qualitatively and quantitatively. The rhizosphere mycoflora was also different in the two varieties tested. Further the inoculated<sup>PT</sup> of plants with Fusarium oxysporum f.sp. ciceri resulted in increase in the mycoflora. In amongst the two cultivars tested, the number of fungi was more in the rhizosphere of JG-62 both in inoculated and uninoculated although more fungi in inoculated rhizosphere. The number of fungi was also more in where oil cakes were used as amendments. The number of fungi in the rhizosphere of inoculated plants of JG-62 with urea was 29, with superphosphate 30, with muriate of potash 27 as against 26 in uninoculated plants; with castor cake 30, with neem cake 32, with mustard cake 34 as against 28 in uninoculated plants; with mahua cake 30 as against 27 in uninoculated plants; with bavistin 24, with vitavax 26, with brassicol 26, with benlate 25, with captan 26, with wettable sulphur 25 in inoculated as against 20, 21, 21, 20, 21 and 21 in uninoculated plants. The number of fungi in the rhizosphere of plants grown in unamended soil was 28 in inoculated as against 26 in uninoculated plants. Almost identical results were obtained with

JG-74 but less number of fungi. The number of fungi was more in the rhizosphere of plants grown in soil amended with fertilizers and oil cakes and less in those treated with fungicides.

As a result of treatment of soil with urea, superphosphate and muriate of potash, almost all the fungi isolated exhibited increased frequency in the rhizosphere of both inoculated and uninoculated plants of JG-62 with respect to control (Table 21) except Syncephalastrum racemosum, Botrytis cinerea, Chaetomium flavum, Aspergillus flavipes and Drechslera hawaiiensis in inoculated and D. hawaiiensis, Alternaria alternata, Fusarium moniliformae, F. oxysporum and Acrophialophora fusipora in uninoculated plants treated with muriate of potash. As a result of treatment with urea the frequency of Rhizopus oryzae increased highest in inoculated and that of Aspergillus fumigatus in uninoculated; with superphosphate Aspergillus niger in inoculated and A. fumigatus in uninoculated and with muriate of potash A. niger in inoculated and Memnoniella echinata and Trichoderma viridae in uninoculated plants.

Saprophytic fungi which exhibited higher frequency in uninoculated plants as compared to inoculated as a result of treatment with urea included A. fumigatus, A. flavus, A. sparsus, A. niger, A. oryzae; with superphosphate A. fumigatus, A. flavus, A. niger, A. sparsus and A. oryzae

and with muriate of potash A. flavus, A. sparsus and T. viridae. Fungi which were present in uninoculated rhizosphere as a result of treatment with urea, superphosphate and muriate of potash included A. fumigatus, A. luchuensis, A. terreus, T. allii, Humicola, A. fuscotra, M. echinata, T. viridae and sterile black mycelium. In addition to this F. moniliiformae with urea and muriate of potash and F. oxysporum with muriate of potash.

With oil cakes not all fungi showed increase in frequency A. fumigatus exhibited highest frequency both in inoculated and uninoculated plants as a result of treatment with castor cake and neem cake; with mustard cake the frequency of A. fumigatus was highest in inoculated and that of A. flavus and sterile black mycelium in uninoculated plants and with mahua cake A. fumigatus in inoculated and A. fumigatus and A. niger in uninoculated plants.

Aspergillus funiculosus, A. luchuensis and sterile black mycelium were present only in the rhizosphere of uninoculated plants treated with castor cake, neem cake, mustard cake and mahua cake. In addition to this T. viridae was also isolated from the rhizosphere of uninoculated plants treated with castor cake F. oxysporum with neem cake and mahua cake and F. moniliiformae with mahua cake.



As a result of treatment with fungicides the frequency of all the fungi both saprophytic and parasitic decreased except A. fumigatus in uninoculated plants treated with different fungicides and that of A. flavus in both inoculated and uninoculated plants treated with bavistin; R. oryzae in inoculated plants treated with bavistin, captan and wettable sulphur and Cladosporium herbarum in inoculated plants treated with bavistin and vitavax. Thus the fungicide had adverse effect on the soil mycoflora both of inoculated and uninoculated plants.

The frequency of pathogenic fungus F. oxysporum f.sp. ciceri with increases urea and superphosphate but muriate of potash, oil cakes and fungicides treatment indicated lower frequency. The frequency ranged from 55 to 85 in fertilizers treated plants and 0 to 25 in oil cakes and fungicides treated plants as against 65 in untreated inoculations. Similarly other pathogenic fungi i.e. Macrophomina phaseolina, A. alternata indicated increase in frequency with fertilizers and decreased with oil cakes and fungicidal treatments. The frequency of former ranged from 20 to 30 in fertilizers treatment and 5 to 15 in oil cake and fungicidal treatment as against 20 in control and that of later ranged from 50 to 30 in fertilizers and 5 to 20 in oil cake and fungicidal treatment as against 30 in control.

Thus oil cake treatment not only brought about decrease in the frequency of pathogenic forms but also increase in the saprophytic forms. Therefore, treatment with oil cakes is more beneficial to the plant and for the disease control.

Almost a similar trend was obtained with JG-74 (Table-23) Cephalosporium aspermum, R. oryzae, Phoma hibernica, A. luchuensis, H. fuscotra, Alternaria humicola and sterile black mycelium were among those saprophytic fungi reported from rhizosphere of uninoculated plants treated with urea, superphosphate and muriate of potash. Frequency of Mucor globosus, A. flavus, A. niger, A. terreus, A. clavatus, Penicillium notatum, Gliocladium roseum, M. echinata was higher in uninoculated plants as compared to inoculated as a result of treatment with urea, superphosphate and muriate of potash Cunninghamella echinulata, A. fumigatus with urea and superphosphate and C. herbarum, D. hawaiiensis and C. lunata with muriate of potash only.

C. aspermum, A. luchuensis, H. fuscotra and A. humicola were isolated from uninoculated plants treated with castor cake neem cake, mustard cake and mahua cake. In addition to this A. sparsus was isolated from the rhizosphere of inoculated plants treated with castor cake, neem cake and mustard cake.

As a result of treatment with fungicides, the frequency of all the fungi both saprophytic and parasitic decreased except A. fumigatus in inoculated and uninoculated plants

treated with different fungicides and A. terreus treated with bavistin, vitavax brassicol and penlate; Rhizopus nodosus in inoculated plants treated with bavistin and captan; Mucor globosus and A. flavus treated with bavistin and vitavax; A. niger treated with vitavax and wettable sulphur; A. clavatus treated with bavistin, vitavax, brassicol and wettable sulphur, T. viridae treated with bavistin brassicol and wettable sulphur and Trichoderma lignorum treated with bavistin, vitavax and captan. The remaining fungi exhibited reduction in frequency as control.

Addition of inorganic fertilizers in soil brought about increase in frequency of almost all the fungi isolated except R. nodosus, C. echinulata, A. alternata as a result of treatment with urea; R. nodosus, M. globosus and A. alternata with superphosphate C. echinulata, C. flavum, P. hibernica, C. lunata and A. alternata in inoculated plants treated with muriate of potash but in uninoculated plants the exception was A. sparsus and A. alternata in all the fertilizers.

Amongst the oil cakes, mustard cake favoured the increase in the frequency of largest number of fungi followed by neem cake, castor cake, and mahua cake in inoculated plants while in uninoculated there has not been a material change, R. nodosus, A. funigatus, A. flavus, A. nigar, A. terreus exhibited increase in frequency in all the oil cakes both in

inoculated and uninoculated plants; C. herbarum in uninoculated treated with castor, neem and mahua cake, A. luceunsi in castor, neem and mustard cake whereas Torulla allii, H. fuscotra, G. roseum, P. notatum in all the oil cake treatments. There was increase in frequency in A. flavus in castor, neem and mustard, Penicillium notatum in castor, neem and mahua and G. roseum in castor and neem cake in inoculated plants.

Fungicidal treatment suppressed the frequency of almost all the fungi both saprophytic and parasitic except a few such as A. fumigatus and A. terreus in some fungicides.

There were some qualitative differences between mycoflora of two varieties, R. nodosus, A. clavatus, P. notatum, M. globosus, G. racemosum and T. lignorum were isolated from inoculated and uninoculated plants, A. terreus, M. echinata, Verticillium glaucum and T. viridae from inoculated and P. hibernica and A. humicola from uninoculated plants of JG-74 and not of JG-62 whereas Acrophialophora fusipora was isolated from both inoculated and uninoculated plants, Mucor racemosus, R. oryzae, Syncephalastrum racemosum, R. oryzae, Syncephalastrum racemosum, M. phasiolina, B. cineria from inoculated and F. oxysporum, F. moniliformae and A. alternata from uninoculated and not from the other cultivar.

A similar trend was observed with rhizoplane fungi with more number of fungi in those inoculated with F. oxysporum f.sp. ciceri (Table 22 and 24). In amongst the two cultivars tested, number of fungi was more in the rhizoplane of cv.JG-62 both in inoculated and uninoculated plants; with urea it was 22 with superphosphate 23 with muriate of potash 21, with castor cake 23, with neem cake 23, with mustard cake 22 with mahua cake 20, with bavistin 16, with brassicol 20, with vitavax 21 with captan 21 against 18 to 21 in uninoculated ones. With benlate and wettable sulphur there has been no difference in the number of fungi in inoculated and uninoculated rhizoplane. In the wettable sulphur the number of fungi was 19 while in benlate was 17, the number of fungi in the rhizoplane of unamended control plants was 21 in inoculated as against 20 in uninoculated plants. With JG-74, the number of fungi was relatively less as compared with JG-62 however fertilizers and oil cakes treatments supported the number of fungi where it was 21 to 24 in inoculated as against 18 to 20 in uninoculated plants. In fungicidal treatment it ranged from 16 to 19 in inoculated as against 17 to 18 in uninoculated plants. The number of fungi control plants was 20 in inoculated as against 18 in uninoculated plants.

Treatment with urea, superphosphate and muriate of potash brought about increase in frequency of almost all the fungi isolated from rhizoplane of JG-62 with respect to control

(Table 22) except C. lunata in uninoculated plants treated with urea; B. cinerea in inoculated plants with superphosphate and C. lunata and A. alternata in both inoculated and uninoculated plants. A. fusipora, R. solani, S. rolfsii, F. oxysporum f.sp. ciceri in inoculated and F. oxysporum, D. hawaiiensis in uninoculated plants with muriate of potash. As a result of treatment with urea the frequency of M. olivaceum was highest in inoculated and that of A. fumigatus in uninoculated plants, with superphosphate R. oryzae in inoculated and A. fumigatus in uninoculated plants.

The frequency of R. oryzae, C. flavum, A. fumigatus, A. flavus, A. niger, A. sparsus was highest in uninoculated as against inoculated plants when treated with urea, superphosphate and muriate of potash. Fungi which are present only in the rhizoplane of uninoculated plants as a result of treatment with urea, superphosphate and muriate of potash included A. terreus, A. lucheunsi, M. echinata, H. fuscotra, T. viridae and sterile black mycelium. In addition to this C. asperum with urea and muriate of potash.

Not all the fungi exhibited increased frequency as a result of oil cake treatment the sterile brown mycelium exhibited highest increase in inoculated and A. fumigatus in uninoculated plants. As a result of treatment with castor cake A. niger in inoculated and A. fumigatus in uninoculated

plants, with neem, mustard and mahua cake C. aspermum, A. terreus, A. luchuensis, M. echinata and H. fuscotra were present only in the rhizoplane of uninoculated plants treated with castor, neem, mustard and mahua cake. In addition to this F. oxysporum was also isolated from rhizoplane of uninoculated plants treated with mustard and mahua cake and T. viridae with mahua cake.

Due to treatment with fungicides, the frequency of all the fungi both saprophytic and parasitic decreased except A. fumigatus in inoculated and uninoculated plants treated with different fungicides and that of A. flavus in inoculated plants treated with bavistin, vitavax and benlate, A. niger with bavistin, vitavax and brassicol, R. oryzae and M. racemosus with vitavax and brassicol and C. echinulata with vitavax only.

The frequency of pathogenic fungus F. oxysporum f.sp. ciceri considerably increased with fertilizers except muriate of potash but oil cakes and fungicides treatment indicated lower frequency. The frequency ranged from 70 to 85 in fertilizers treated plants and 0 to 25 in oil cakes and fungicides treated plants as against 60 in untreated inoculation. Similarly, other pathogenic fungi i.e. A. alternata indicated increase in frequency with fertilizers and decreased with oil cake and fungicidal treatments. The frequency ranged from 40 to 50 in urea and superphosphate and 5 to 20 in muriate

of potash, oil cakes and fungicidal treatment as against 30 in control.

C. aspermum, H. fuscotra and A. humicola among those saprophytic fungi reported from rhizoplane of uninoculated plants of JG-74 treated with urea, superphosphate and muriate of potash. In addition to this R. oryzae, A. sparsus from rhizoplane of uninoculated plants treated with urea and superphosphate. Frequency of A. flavus, A. niger, A. terreus, A. clavatus, M. echinata and T. viridae was higher in uninoculated plants as compared to inoculated as a result of treatment with urea, superphosphate and muriate of potash, A. fumigatus with urea and superphosphate and C. herbarum with urea. C. aspermum, A. humicola were isolated from uninoculated plants treated with castor cake, neem cake, mustard cake and mahua cake. As a result of treatment with fungicides the frequency of all the fungi both saprophytic and parasitic decreased except A. fumigatus in inoculated and uninoculated plants treated with different fungicides. R. nodosus in inoculated plants with bavistin, vitavax and captan; M. globosus with bavistin and vitavax; A. flavus with bavistin; A. niger with bavistin, brassicol, captan and wettable sulphur; A. clavatus with bavistin, vitavax, brassicol, benlate and wettable sulphur and T. viridae with bavistin, vitavax, brassicol and wettable sulphure. All the remaining fungi exhibited reduction in frequency as compared

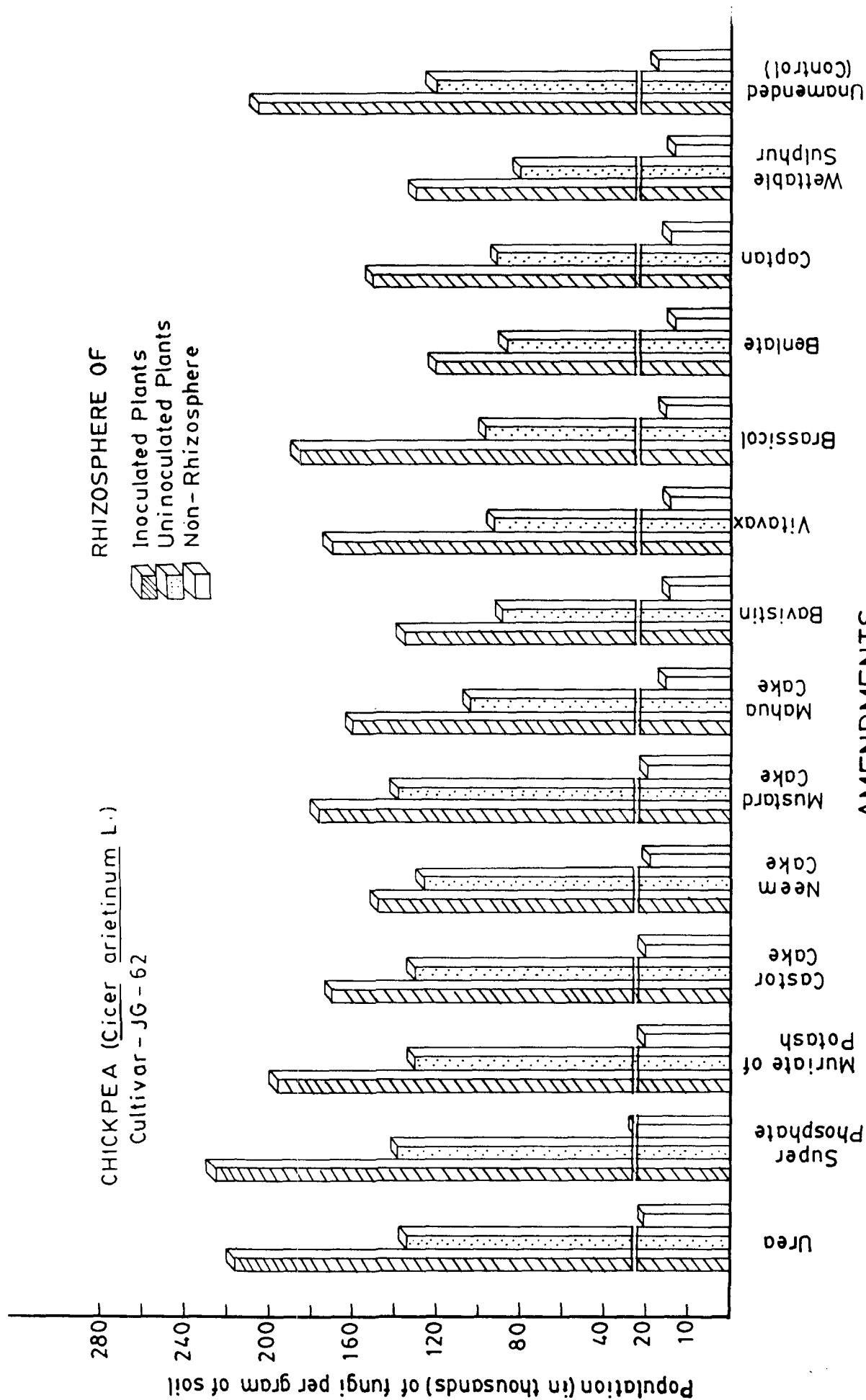


to control. A similar trend was observed in the reduction in the frequency of pathogenic fungi.

The two cultivars also exhibited differences in rhizoplane. M. globosus, G. roseum and A. humicola were isolated from JG-74 while A. luchuensis, B. cinerea, A. fusipora and F. oxysporum from JG-62.

It is interesting to note that the frequency of saprophytic fungi such as Aspergillus species was more in the rhizoplane of JG-74 both in inoculated and uninoculated plants as a result of treatment with different fertilizers, oil cakes and fungicides. On the other hand, the frequency of pathogenic fungi was more in the rhizoplane of JG-62 due to various treatments. It is likely that higher frequency of saprophytic fungi in JG-74 is a contributory factor towards less development of disease on JG-74.

FIG. 5. Population of fungi in the non-rhizosphere, rhizosphere of uninoculated and inoculated plants of chickpea cultivar JG-62 (with Fusarium oxysporum f.sp. ciceri) treated with different soil amendments.



AMENDMENTS

Fig.5

FIG. 6. Population of fungi in the non-rhizosphere, rhizosphere of uninoculated and inoculated plants of chickpea cultivar JG-74 (with Fusarium oxysporum f.sp. ciceri) treated with different soil amendments.

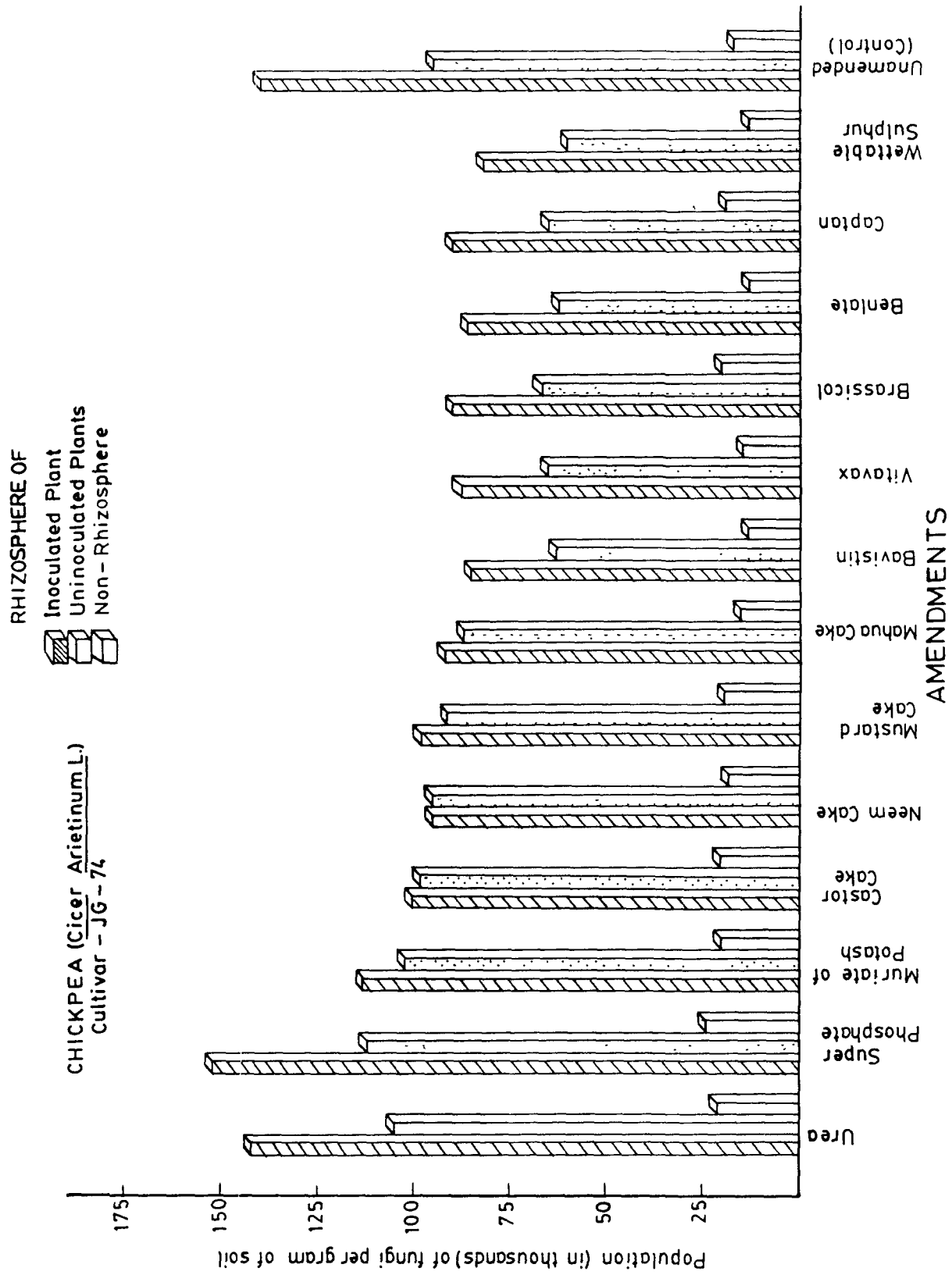


Fig. 6

TABLE - 20: Population of fungi (per gram of soil) in the non-rhizosphere and rhizosphere of chickpea plants (cultivar JG-62 and JG-74) uninoculated and inoculated with Fusarium oxysporum f.sp. ciceri under different soil amendments

Amendments	Non-rhizosphere	RHIZOSPHERE					
		JG-62			JG-74		
		Inoculated Plant	R:S Ratio	Uninoculated Plant	R:S Ratio	Inoculated Plant	R:S Ratio
Control	17.0	205.0	12.05	120.0	7.05	140.0	8.23
Urea	21.5	216.5	10.06	134.0	6.23	146.0	6.79
Superphosphate	24.0	225.0	9.37	138.5	5.77	142.0	5.91
Muriate of Potash	20.5	195.0	9.51	131.5	6.41	113.0	5.50
Castor Cake	20.0	170.0	8.50	129.0	6.45	100.0	5.00
Neem Cake	18.5	148.0	8.00	126.0	6.81	95.0	5.13
Mustard Cake	19.0	176.0	9.26	128.5	6.76	98.0	5.15
Mahua Cake	14.5	160.0	11.03	104.5	7.17	92.5	6.37
Bavisting	13.5	135.0	10.00	88.5	6.55	85.5	6.33
Vitavax	14.0	170.0	12.14	92.0	6.57	87.5	6.25
Brassiccol	15.0	185.0	12.33	95.5	6.36	90.0	6.00
Benlate	13.0	120.0	9.23	85.0	6.53	86.0	6.60
Captan	14.0	150.0	10.71	90.0	6.42	90.0	6.42
Wettable Sulphur	12.5	130.0	10.40	80.0	6.40	82.0	6.56
L.S.D. at 5%	1.3367147	0.955214		1.2423029		1.0873148	
L.S.D. at 1%	1.816875	1.2983353		1.6885497		1.477883	

statistical analysis

TABLE - 21: Frequency (percentage) of fungi in the rhizosphere of chickpea plants cultivated under different soil amendments, and inoculated with Fusarium oxysporum f.sp. ciceri

Fungi isolated	1		2		3		4		5		6		7		8		9		10		11		12		13		14			
	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP		
<u>Rhizopus oryzae</u>	15	30	35	40	30	40	25	35	30	35	25	30	20	30	20	25	15	20	15	20	15	15	10	15	20	25	10	20		
<u>Mucor racemosus</u>	10	-	20	-	25	-	15	-	5	5	10	5	10	10	5	5	-	10	-	5	-	5	5	-	5	-	20	-		
<u>Cunninghamella echinulata</u>	20	-	25	-	30	-	25	-	10	10	10	10	15	10	15	15	-	25	-	20	-	20	-	5	-	15	-	10	-	
<u>Mortierella alpina</u>	15	-	25	-	35	-	20	-	15	-	10	-	10	-	10	5	-	5	-	5	-	5	-	5	-	10	-	5	-	
<u>Cephalosporium asperum</u>	20	15	30	25	35	30	25	20	10	10	10	10	15	10	10	5	-	5	-	5	-	10	-	5	-	15	-	10	-	
<u>Monilia humicola</u>	30	25	50	40	45	40	40	35	30	30	30	25	30	25	15	10	15	15	20	10	20	15	15	10	20	15	15	10		
<u>Syncephalastrum racemosum</u>	20	-	40	-	35	-	15	-	20	-	15	-	15	-	15	-	10	-	15	-	10	-	10	-	10	-	10	-		
<u>Chaetomium flavum</u>	15	20	40	30	50	35	25	25	15	20	20	20	15	10	20	15	15	10	10	10	20	15	10	10	15	10	10	10		
<u>Macrophomina phaseolina</u>	20	-	30	-	20	-	25	-	15	-	5	-	10	-	-	-	-	-	5	-	5	-	5	-	10	-	10	-		
<u>Aspergillus fumigatus</u>	35	25	50	70	55	70	65	45	55	75	50	80	40	40	50	45	35	40	35	45	40	30	30	35	25	35	20	25		
<u>A. flavus</u>	20	25	40	50	50	55	35	45	30	45	30	40	25	50	30	40	30	30	40	15	15	20	20	15	10	20	10	15		
<u>A. niger</u>	15	30	30	40	40	45	30	35	20	50	25	45	20	40	30	45	20	20	15	25	15	25	10	15	25	25	20	15		
<u>A. funiculosus</u>	-	20	-	25	-	30	-	25	-	25	-	25	-	20	-	20	-	15	-	10	-	10	-	10	-	15	-	10	-	
<u>A. terreus</u>	-	20	-	40	-	45	-	35	10	40	20	35	15	30	15	35	-	15	-	10	-	15	-	10	-	10	-	10	-	
<u>A. luchuensis</u>	-	10	-	15	-	20	-	15	-	25	-	25	-	20	-	25	-	-	-	-	-	-	-	-	-	-	-	5	-	
<u>A. sparsus</u>	15	20	25	35	30	30	20	35	25	30	20	30	20	25	20	25	10	10	10	15	10	10	10	10	5	10	5	15		
<u>A. candidus</u>	15	-	25	-	30	-	20	-	15	-	20	-	20	-	20	-	10	-	10	-	10	-	5	-	10	-	5	-		
<u>A. flavipes</u>	20	15	30	25	35	30	20	20	30	20	25	20	20	20	30	25	10	-	5	-	5	-	5	-	10	10	5	-		
<u>Monosporium olivaceum</u>	1	-	40	-	-	-	10	-	10	-	1	-	5	-	10	-	-	-	-	-	-	-	-	-	-	-	-	-	-	
<u>Botrytis cinerea</u>	20	-	40	-	-	-	15	-	-	-	10	-	5	-	10	-	10	-	15	-	15	-	10	-	10	-	10	-	-	
<u>Torula alli</u>	-	0	-	10	-	10	-	35	10	25	10	30	10	30	10	25	-	10	-	15	-	15	-	10	-	15	-	10	-	
<u>Humicola fuscolata</u>	-	20	-	10	-	10	-	30	10	20	10	20	15	35	10	25	-	15	-	20	-	15	-	15	-	20	-	15	-	
<u>Memnoniella echinata</u>	-	25	-	10	-	15	-	50	20	30	10	30	10	40	15	25	10	10	-	10	-	10	-	10	-	15	-	15	-	
<u>Gladosporium barbarum</u>	30	25	25	10	10	10	45	35	20	30	10	20	15	20	15	20	20	20	40	20	20	15	15	20	20	20	15	10		
<u>Drechslera nivalensis</u>	25	20	10	10	10	10	25	20	10	10	10	10	10	10	15	10	15	10	15	10	15	20	15	15	15	15	10	10		
<u>Curvularia lunata</u>	25	20	10	10	10	10	30	25	15	10	10	10	10	10	15	15	10	15	10	15	10	15	10	10	10	10	10	10		
<u>Alternaria alternata</u>	30	20	20	10	10	10	40	30	15	10	10	10	10	10	15	10	10	10	15	15	15	15	15	10	10	10	10	10		
<u>Fusarium moniliformae</u>	-	20	-	10	10	10	10	20	20	20	15	10	10	10	10	10	10	10	10	10	10	10	15	10	10	15	15	10		
<u>F. oxysporum</u>	-	30	10	10	10	10	10	20	20	20	15	10	10	10	10	10	10	10	10	10	10	10	15	10	10	15	15	10		
<u>F. oxysporum f.sp. ciceri</u>	65	-	70	-	10	-	55	-	15	-	-	-	25	-	-	-	-	-	15	-	20	-	-	-	20	-	10	-		
<u>Trichosporium fuscum</u>	20	10	30	20	30	20	25	15	15	5	20	10	25	15	10	-	5	-	5	-	10	-	-	-	5	-	-	-	-	
<u>Epicarpha silvatica</u>	25	15	35	25	40	25	30	20	10	10	15	15	30	20	5	-	10	-	10	-	15	-	5	-	-	-	-	-	-	
<u>Trichoderma viridae</u>	-	20	-	35	-	45	20	40	-	20	10	15	10	20	5	15	-	15	-	10	-	20	-	15	-	10	-	10	-	
<u>Acrophalophora fusipora</u>	15	10	30	20	35	20	20	10	-	5	10	5	5	10	10	5	5	5	5	5	5	5	5	5	5	5	5	-	-	
<u>Sclerotium rolfsii</u>	10	-	20	-	15	-	-	-	-	-	-	-	5	-	-	-	-	-	-	-	-	-	-	-	-	-	5	-	-	
<u>Rhizoctonia solani</u>	15	-	20	-	10	-	-	-	-	-	5	-	5	-	-	-	-	-	-	-	-	-	5	-	5	-	5	-	-	
<u>Sterile brown mycelium</u>	25	-	40	-	45	-	35	-	35	-	35	-	45	-	30	-	10	-	5	-	15	-	15	-	25	-	15	-	-	
<u>Sterile black mycelium</u>	-	30	-	55	-	50	-	45	-	40	-	40	-	50	-	40	-	15	-	15	-	20	-	20	20	-	20	-	10	-

1. Control, 2. Urea, 3. Superphosphate, 4. Furiate of potash, 5. Castor cake, 6. Neem cake, 7. Mustard cake, 8. Mahua cake, 9. Bavistin, 10. Vitavax, 11. Brassicol, 12. Benlate, 13. Captan, 14. Wettable sulfur  
IP = Inoculated plants, UP = Uninoculated plants

Urea. 2. Urea. 3. Superphosphate. 4. Muriate of potash. 5. Castor cake. 6. Neem cake. 7. Mustard cake. 8. Mahua cake. 9. Bavistar, lc. V. vavax.

UP = Uninoculated plants

Childs  
1907



TABLE - 23: Frequency (percentage) of fungi in the rhizosphere of chickpea plants cultivar J0-74 uninoculated and inoculated with *Fusarium oxysporum* f.sp. *clavari* under different soil amendments

	1	2	3	4	5	6	7	8	9	10	11	12	13	14												
	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP	IP	UP												
<i>Phaeoacremonium</i>	-	25	-	35	-	40	-	30	10	30	20	25	10	30	20	40	-	10	-	20	-	20	-	15	-	15
<i>Phaeoacremonium</i>	20	15	15	20	15	25	30	20	35	30	30	20	25	15	20	20	20	5	15	10	15	15	10	10	15	10
<i>Phaeoacremonium</i>	25	-	20	-	20	-	25	-	15	10	10	-	10	-	15	-	10	-	15	-	20	-	10	-	15	-
<i>Phaeoacremonium</i>	15	10	20	20	10	20	10	15	5	5	10	5	5	5	10	10	15	-	20	-	10	-	5	-	5	-
<i>Phaeoacremonium</i>	10	-	15	-	20	-	15	-	10	-	5	-	5	-	5	-	5	-	10	-	5	-	-	-	-	-
<i>Phaeoacremonium</i>	20	20	30	25	35	25	25	25	10	15	10	10	5	10	15	15	10	10	15	15	10	15	10	10	10	10
<i>Phaeoacremonium</i>	-	30	-	35	-	40	-	35	-	20	-	10	-	10	-	15	-	15	-	25	-	20	-	10	-	20
<i>Phaeoacremonium</i>	20	-	30	-	30	-	15	-	15	-	15	-	10	-	15	-	15	-	10	-	10	-	10	-	10	-
<i>Phaeoacremonium</i>	-	15	-	30	-	25	10	30	5	30	15	30	10	15	5	20	-	10	-	15	-	10	-	10	-	10
<i>Phaeoacremonium</i>	30	25	60	80	55	75	70	40	60	85	45	60	55	70	60	75	35	30	35	45	30	40	30	30	30	30
<i>Phaeoacremonium</i>	15	25	25	35	30	35	20	30	20	40	25	45	20	35	30	40	20	15	15	20	15	10	10	10	10	10
<i>Phaeoacremonium</i>	20	35	55	65	60	70	65	75	65	65	70	70	40	45	50	60	20	20	25	30	20	30	15	10	10	10
<i>Phaeoacremonium</i>	25	-	30	-	35	-	30	-	30	20	30	-	35	-	25	-	5	-	15	-	10	-	10	-	15	-
<i>Phaeoacremonium</i>	20	35	50	70	45	65	55	75	60	80	55	75	45	70	40	65	20	40	30	40	15	45	20	10	15	15
<i>Phaeoacremonium</i>	-	20	-	25	-	30	15	30	-	25	-	25	-	25	-	20	-	10	-	5	-	15	-	10	-	10
<i>Phaeoacremonium</i>	15	-	20	-	20	-	15	-	-	-	-	-	15	-	15	-	20	-	15	-	10	-	10	-	10	-
<i>Phaeoacremonium</i>	10	25	20	40	20	50	25	45	10	40	15	50	15	40	10	40	15	15	15	20	15	20	10	10	15	20
<i>Phaeoacremonium</i>	-	15	-	20	-	25	-	20	-	25	-	25	-	25	-	10	30	-	15	-	10	-	10	-	10	-
<i>Phaeoacremonium</i>	20	35	35	40	30	45	25	40	30	45	30	40	20	45	25	40	10	15	15	30	10	25	10	20	15	25
<i>Phaeoacremonium</i>	30	40	40	50	35	55	35	45	45	50	35	50	25	60	30	50	10	20	20	30	15	25	25	35	20	30
<i>Phaeoacremonium</i>	20	-	25	-	25	-	25	-	15	-	10	-	5	-	10	-	-	5	-	10	-	10	-	10	-	10
<i>Phaeoacremonium</i>	-	25	-	35	-	40	-	30	15	30	10	35	-	40	-	35	-	10	-	20	-	15	-	10	-	15
<i>Phaeoacremonium</i>	25	-	30	-	35	-	30	-	35	-	30	-	35	-	35	-	15	-	20	-	20	-	20	-	25	-
<i>Phaeoacremonium</i>	-	20	-	25	-	30	-	30	-	30	-	35	-	30	-	35	-	10	-	15	-	15	-	10	-	10
<i>Phaeoacremonium</i>	15	35	30	65	35	55	30	70	15	45	20	40	15	45	20	40	10	30	10	20	15	20	10	25	10	25
<i>Phaeoacremonium</i>	35	30	45	40	45	45	15	40	30	15	25	20	20	15	20	20	25	10	20	25	25	20	15	10	25	15
<i>Phaeoacremonium</i>	35	30	40	35	55	60	25	35	20	20	30	25	30	25	25	20	15	15	15	15	15	15	10	30	25	10
<i>Phaeoacremonium</i>	40	35	60	50	50	55	20	40	20	20	30	20	25	20	20	20	20	25	20	25	20	25	20	25	20	20
<i>Phaeoacremonium</i>	20	-	20	-	25	-	10	-	15	-	10	-	5	-	10	-	15	-	10	-	10	-	5	-	10	-
<i>Phaeoacremonium</i>	-	15	-	20	-	25	-	20	-	15	-	20	-	20	-	25	-	-	-	-	-	5	-	10	-	-
<i>Phaeoacremonium</i>	40	40	45	-	50	-	40	-	-	-	5	-	20	-	-	-	-	-	20	-	-	-	-	-	-	-
<i>Phaeoacremonium</i>	10	30	20	35	15	45	25	40	15	40	20	40	15	40	10	25	15	15	10	10	10	10	10	10	10	10
<i>Phaeoacremonium</i>	10	15	15	25	15	25	25	25	15	25	20	25	20	30	15	30	15	10	15	10	15	10	10	10	10	10
<i>Phaeoacremonium</i>	10	-	15	-	25	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-	-
<i>Phaeoacremonium</i>	25	-	40	-	35	-	40	-	35	-	40	-	50	-	40	-	15	-	-	-	-	-	-	-	-	-
<i>Phaeoacremonium</i>	-	20	-	30	-	40	-	45	-	35	-	40	-	45	-	45	-	1	-	1	-	1	-	10	-	-

1 Control, 2. Urea, 3. Superphosphate, 4. Muriate of potash, 5. Castor cake, 6. Neem cake, 7. Mustard cake, 8. Brackish water, 9. Brackish water, 10. Vitex, 11. Brackish water, 12. Benlate, 13. Captan, 14. Wettable sulphur; IP = Inoculated plants, UP = Uninoculated plants.

Handwritten signature and date: 10/10/10

1. ATTENDANCE  
11

in the range of 0.10

of two or three inches.

12. COURT, the

1

ricer[illegible]

Control, 2. Urea, 3. Superphosphate, 4. Muriate of potash, 5. Castor cake, 6. leen cake, 7. mustard cake, 8. Mahua cake, 9. 3 v l t', 10. 1/10 x,

- Jassicol, 12. Penlate, 13. Carton, 14. Wetable sulphur

recultivated plants,  $\chi^2 = 11.90$  1 ted plants

phur  
stat. needed  
is

## CHAPTER - IV

### DISCUSSION

Chickpea (Cicer arietinum L.), an important crop of dryland farming is a good source of protein. Amongst, different diseases which damage the crop, wilt of chickpea caused by Fusarium oxysporum f.sp. ciceri is of (no less) importance. Being <sup>the fungus</sup> a soil borne, it influences the soil microflora and also is influenced by the soil microflora. Amongst the different soil microflora, some are advantageous and others disadvantageous to the organisms causing the disease. Those disadvantageous to causal organism help the plant to overcome the pathogenic effect and have been exploited for biological control of the disease. The aim of the present studies has been to determine the soil mycoflora of chickpea infected with F. oxysporum f.sp. ciceri and to determine the changes in the soil mycoflora as a result of foliar sprays with growth regulators, fertilizers and pesticides and soil amendment with oil cakes, fertilizers and fungicides so as to understand the dynamics of soil mycoflora.

The population of fungi has been high in the rhizosphere of both inoculated and uninoculated plants as compared to

non-rhizosphere. This is understandable as the rhizosphere zone is rich in amino acids, carbohydrates and growth promoting substances in the form of root exudation<sup>7</sup> which have already been reported to have stimulating effect on fungi. These findings are thus in agreement with earlier studies (Starkey, 1929, 1931 and 1958; Timonin, 1940a,b; Lochhead, 1959; Maliszewska and Moreau, 1959; Reddy, 1959; Ivarson and Katznelson, 1960; Rauatt and Katznelson, 1961; Zagallo and Bollen, 1962; Rangaswamy and Vasantharajan, 1962; Kovira, 1963; Singh, 1971; Rai and Upadhyay, 1980 and Tandon and Tiwari (1982). Moreover, the fungal population has been high in the rhizosphere of plants inoculated with wilt causing organism as compared to uninoculated ones throughout the studies. In the diseased plants there has been greater metabolic activity leading to more root excretions, which might be one of the reasons for more activity of fungi (Agnihotrudu, 1957 and 1959; Timonin, 1966; Wood, 1967; Mathur and Chauhan, 1972; Babushkina., 1973; Rai and Upadhyay, 1980; Kumar and Balasubramanian, 1981; Satyaprasad, 1982; Wadhwani and Mehrotra, 1982; Vesely, 1985 and Suteri, 1986). This is further supported by the fact that the concentrations of total free amino acids, phenols, O-dihydroxy phenols and sugars has been higher in the roots of inoculated than to uninoculated plants. These compounds might have leached out in the soil thus providing more stimulation of fungi (Wood, 1967; Mathur and Chauhan, 1972; Srivastava and Mishra, 1972;

Khan et al., 1973; Babushkina, 1973; Rai and Upadhyay, 1980; Ashour et al., 1980; Wadhwani and Mehrotra, 1982; Kishan et al., 1982; Ansari and Prakash, 1985 and Suteri, 1986). Satyaprasad and Rama Rao (1983) also concluded that amino acids and sugars in the root exudates exert a marked influence on germination of conidia and chlamydospore<sup>s</sup> of F. oxysporum f.sp. ciceri in the root region leading to root infection.

The rhizosphere and rhizoplane mycoflora of different cultivars of chickpea inoculated with F. oxysporum f.sp. ciceri exhibit differences both qualitatively and quantitatively. Cultivar H-208 harboured <sup>the</sup> highest number of fungi in the rhizosphere, while H-208 and JG-62 <sup>with the highest number</sup> in the rhizoplane. The fungus population has also been high in the cultivar H-208 and least in JG-74, in both inoculated and uninoculated plants. Amongst different fungi isolated the frequency of Aspergillus niger has been high in the rhizosphere and rhizoplane of uninoculated plants of all the cultivars while that of F. oxysporum f.sp. ciceri in the rhizosphere and rhizoplane of those cultivars which happen to be susceptible to the disease. However in BG-315, BG-212 and JG-74, which (otherwise) <sup>are</sup> less prone to disease, Aspergillus fumigatus and A. flavus dominated. In all the cultivars, the frequency of parasitic forms has been high in those inoculated with F. oxysporum f.sp. ciceri and those of saprophytic fungi

in uninoculated plants (Table - 4,5). This difference in the fungal flora and population in the rhizosphere and rhizoplane of various cultivars could be due to variations in the root exudates. These results thus are in conformity with those of Subbarao and Bailey (1961), Parkinson (1967), Gujrati (1969), Jalali and Suryanarayana (1972), Sullia (1973), Srivastave (1974), Rataj Guranowsky (1981) and Tandon and Tiwari (1982). The frequency of antagonistic fungi such as Trichoderma viridae and Aspergillus terreus has been high in rhizosphere<sup>2</sup> of those cultivars where the frequency of F. oxysporum f.sp. ciceri has been low and which are less prone to disease. This possibly explains in part the poor development of F. oxysporum f.sp. ciceri in such cultivars.

It is interesting to note that the cultivars which harbour more fungi and have higher frequency of F. oxysporum f.sp. ciceri, both in the rhizosphere and rhizoplane, have usually low concentrations of phenols, <sup>only</sup> 0-dihydroxy phenols but higher concentration of sugars (Table -2). The reverse was true with cultivars which harbour lower number of fungi together with lower frequency of F. oxysporum f.sp. ciceri. The role of phenols and 0-dihydroxy phenols in the resistance of plants against disease has already been emphasized (Sharma et al., 1982 and Mikzak et al., 1985a). Alanine, <sup>a</sup>Aspartic acid, <sup>a</sup>Glutamic acid and <sup>a</sup>Glycine have been detected in the root

extracts of all the cultivars while Asparagine, Lysine and Histidine in JG-62, H-208, JG-309 and BG-212; Serine in H-208, BG-309, JG-313, BG-212 and JG-74; Phenylalanine in JG-62, H-208, BG-309, BG-212 and JG-74; Tryptophane and Valine in JG-62, H-208 and JG-309; Tyrosine in JG-315, BG-212, and JG-74; Cysteine in BG-212 and JG-74 and Lucine in JG-315. The number of amino acids however, varies from 8-12 in different cultivars, the highest being with H-208 and BG-309 followed by JG-62, JG-315, BG-212 and JG-74. There has been, however, no difference in the number of amino acids in inoculated and uninoculated plants of different cultivars. The number of amino acids have been relatively high in the cultivars susceptible to wilt causing organism i.e. JG-62, H-208 and BG-309 as compared to BG-212 and JG-74 which are less damaged. The amino acid and sugar composition of the root exudates is likely to affect the C/N ratios, which in turn affect the growth of fungi in the root region (Cook and Schroth, 1965). It is likely that more amino acids together with high concentration<sup>s</sup> of sugar might be favouring the growth of F. oxysporum f.sp. ciceri in the soil and inside the roots. Subbarao and Bailey (1961) also reported that<sup>a</sup> susceptible variety of tomato to Verticillium wilt contained more amino acids in the root exudates than the resistant variety. Satyaprasad and Rama Rao (1983) also found more amino acids in the root exudates of susceptible in comparison to resistant variety of chickpea. It has been

suggested that higher number of amino acids possibly contribute more soluble nitrogenous material to the rooting medium and thus help the pathogen to effectively colonize. In the present studies cysteine has been detected in cv. BG-212 and JG-74 which are less damaged by the wilt causing organisms. This might be <sup>a</sup>contribution towards the resistance of plants against the disease. Similar correlation of resistance <sup>to amino acid content</sup> have been made in cotton to Xanthomonas campestris pv. malvacearum (Vohra and Chand, 1971) and greengram to Xanthomonas campestris pv. phaseoli (Marimuthu, 1983). ~~Lysine, Valine, Asparagine and Histidine~~, which were detected only in the roots of the susceptible variety, are already known for their stimulatory effect on the growth of fungi (Leonian and Lilly, 1940; Lilly and <sup>W</sup>Brnett, 1951; Cochrane, 1958 and Schroth and Snyder, 1961).

The increase <sup>d</sup>concentration of amino acids in infected plants could in part be due to degradation of protein into amino acids due to production of toxins by the pathogen, and/or activation of enzymes involved in the amino acid and amide synthesis, and/or inhibition of enzymes by the pathogen which are involved in the protein synthesis (Goodman et al., 1967; Tandon, 1970 and Jalali and Suryanarayana, 1972).

*Review Not a full*  
In the studies dealing with the age of plants on rhizosphere and rhizoplane mycoflora. The population of fungi increased with the increase in age of plant upto





the other hand, the frequency of saprophytic fungi was higher in uninoculated plants. This indicates that in the rhizosphere of inoculated plants, parasitic fungi, particularly F. oxysporum f.sp. ciceri dominate.<sup>The</sup> Reverse was, however, true with uninoculated plants. Similar results were obtained by Vesely (1985) with sugarbeet in relation to dampin-off fungus.

As a result of inoculation, differences have been observed in the rhizosphere and rhizoplane mycoflora of diseased and healthy plants at different stages of plant growth. Similar results were reported by Parlik, 1950; Timonin, 1966; Hong, 1969; Rai and Upadhyay, 1980 and Vesely, 1985. It has been reported that root exudates in the form of amino acids and carbohydrates have a distinct selective action on the rhizosphere microorganisms resulting in the stimulation of certain groups and suppression of the others (Lochhead, 1940; Gujrati, 1969 and Tandon and Tiwari, 1982). In the beginning, infection by the pathogen induces changes in chemical composition of the host which might be advantageous to some fungi and disadvantageous to others. Higher concentration<sup>S</sup> of amino acids, phenols, O-dihydroxy phenols and sugars have been found in the inoculated plant roots in comparison to uninoculated counterparts. These results are in conformity with those of Wood (1967), Jalali and Suryanarayana (1972), Singh

et al. (1978) and Suteri (1986). Karimbaeva and Sizova (1977) reported a direct relationship of concentration of root exudates on growth of fungi.

Certain forms of fungi are recorded throughout the growth period of plants; however, others are restricted to a particular stage of growth. By and large, Aspergillia spp. constitute dominant mycoflora throughout the growth period in uninoculated plants but in the inoculated plants, F. oxysporum f.sp. ciceri and other parasitic fungi constitute the dominant mycoflora throughout the growth of plants. This is in conformity with Mathur and Chauhan (1972) and Satyaprasad and Rama Rao (1983). The variation in the rhizosphere and rhizoplane mycoflora due to age of plant may probably be due to qualitative and quantitative changes in the root exudates at various growth stages as reported by Scroth and Hildbrend (1964), Katznelson (1965), Rovira (1965) and Vesely (1985). In the present studies there has been an increase in the concentration of total free amino acids, phenols, O-dihydroxy phenols and sugars in the roots (of inoculated and uninoculated) with the increase in the age of plants upto 90 days followed by a decline.

A slight decline in the population of fungi after fruiting stage may be attributed due to moisture stress as in this crop only one irrigation is done during plant growth, or in part may be due to seasonal variation in temperature

and moisture. Gangawane and Deshpande (1977) also explained that qualitative and quantitative differences in the fungal flora could be due to a response to seasonal variations. Gujrati (1968) showed that a change in the moisture content had a pronounced effect on the fungal flora of chickpea and lentil. The activity of F. orthoceras var. ciceri has been correlated with moisture content of soil (Chauhan, 1963). Bissett and Parkinson (1979) also pointed out that temperature, moisture, available K, and soil pH are the most important abiotic variables influencing the distribution and composition of soil mycoflora.

Sprays with growth promoting substances have been found to increase the population<sup>S</sup> of fungi in the rhizosphere<sup>S</sup> of both the cultivars JG-62 and JG-74 except maleic hydrazide where there has been some reduction (Table - 10,11). Similar observations regarding changes in rhizosphere population<sup>S</sup> of fungi due to foliar sprays with growth promoting substances have been reported by Vransy et al. (1962), Ray and Dwivedi (1967), Sullia (1968), Misra and Gujrati (1969), Singh (1970), Gupta (1971), Dwivedi and Singh (1971) and Singh (1981). It appears that sprays with chemicals might influence the metabolic activity in plants (Singh and Murty, 1987) which in turn influence the root exudation. The<sup>+</sup> root exudates influencing the<sup>F</sup> rhizosphere population<sup>S</sup> has already been documented (Vransy et al., 1962, Agnihotri, 1964 and Singh, 1981)

*80% of spray is absorbed by the plant*

The spray<sup>ing</sup> of plants with indole acetic acid, indole butyric acid, thio-indole butyric acid and gibberellic acid results in increase<sup>s</sup> in the frequency of fungi; however, with maleic hydrazide there has been a decrease of majority of fungi (Tables - 12, 14, 16 and 18). There has been more increase in the frequency of saprophytic fungi more in the rhizosphere of uninoculated<sup>than in inoculated</sup> plants. With maleic hydrazide spray, there has been some increase in the frequency of A. fumigatus in inoculated plants, A. niger in uninoculated plants of both ~~the~~ cultivars; Trichoderma lignorum and Fusarium oxysporum in uninoculated plants of JG-74 and JG-62 respectively. The differences could be due to changes in the root exudates of the plants. In plants sprayed with growth promoting substances, the growth promoters are absorbed by the leaves and undergo metabolic changes in the tissue leading to activation of various enzymes such as peroxidases etc. (Sembdner et al., 1980). The metabolic products are leached out of the roots. It is likely that the metabolic products of maleic hydrazide leached out are toxic for most of the fungi in the rhizosphere (Dwivedi and Singh, 1971). The increase of certain saprophytic forms such as A. niger, T. lignorum in the rhizosphere of plants treated with maleic hydrazide is of some importance which could be used for disease management (Mathur and Chauhan, 1972; Gokulapalan and Nair, 1984; Venkatasubbaiah and Safeulla, 1984 and

Zozzerini and Tosi, 1985).

Results on the effect of foliar sprays with fertilizers indicate that, by and large, sprays with urea exhibit stimulatory effect on the rhizosphere population of fungi both in inoculated and uninoculated plants while muriate of potash <sup>has</sup> an inhibitory effect (Tables - 10,11). These results are in agreement with Agnihotri (1964), Ramachandra Reddy (1959 and 1968), Dwivedi and Singh (1971) and Rao and Raja (1978) who reported an increase in rhizosphere population of fungi with sprays of urea but are slightly in disagreement with Vransy (1963 and 1972) and Annapurna and Rao (1983) who point out decrease in the number of fungi as a result of spray.

Significant effect of spray <sup>in</sup> with urea on frequency of rhizosphere fungi has been observed only after <sup>the</sup> 2nd and 3rd sprays in both <sup>of</sup> the cultivars (Tables - 12,14,16 and 18). The urea when sprayed is probably absorbed by the leaves and is transformed (Throne, 1954 and Boynton, 1954) into more amino acids such as glutamine, glutamic acid and  $\alpha$ -aminobutyric acid, which exude out from the roots (Agnihotri, 1964 and Steward and Durzan, 1965). These might be providing <sup>a</sup> stimulatory effect on fungi (Leonian and Lilly, 1940, Lilly and Barnett, 1951, Cochrane, 1958 and Schroth and Snyder, 1961).

disc junction sentence  
P. 112

With muriate of potash, although overall decrease has been observed in mycoflora but while comparing the two groups of fungi i.e., parasitic and saprophytic, the saprophytic fungi indicate an increase while that of parasitic fungi a decrease. High amount<sup>S</sup><sub>n</sub> of potassium in soil <sup>is</sup> believed to reduce infection with several soil borne fungi (Lewis, 1979 and Huber, 1979). The increase in potassium levels might increase the vigour of plants thus resulting in root exudates unfavourable to parasitic fungi of rhizosphere.

It is interesting to note that in the rhizosphere of inoculated plants of JG-74, a cultivar <sup>with moderate resistance?</sup> ~~(relatively less susceptible)~~ to F. oxysporum f.sp. ciceri, there has been a significant increase in frequency <sup>of</sup> I. viridae with spray<sup>S</sup><sub>n</sub> of potash, (the fungus known for antibiosis) (Gokulapalan and Nair, 1984) <sup>giving li.</sup> ~~might in part be responsible for reduction in frequency of parasitic forms in these studies.~~ This <sup>may</sup> ~~can~~ be exploited for disease control (Martinez Vicra and Garcia Gomez, 1984).

P. 112  
No significant  
variation  
observed.

Inhibition in the population of rhizosphere fungi has also been observed with sprays of bavistin, vitavax, brassicol, benlate, captan and wettable sulphur both in inoculated and uninoculated plants in both the cultivars JG-62 and JG-74 (Tables-10,11). This kind of inhibition in the rhizosphere mycoflora with fungicidal sprays has also been observed by Helleck and Cochrane (1950), Sullia (1969), Srivastava and Mishra (1971), Ranga Rao et al. (1972), Balasubramanian and

Rangaswami (1973), Wainwright and Sowden (1977), Srivastava and Dayal (1981), Gunaseka and Rao (1982) and Abdul Kader et al. (1983). The frequency of majority of the parasitic fungi including F. oxysporum f.sp. ciceri decrease due to foliar sprays with fungicides in the rhizosphere and rhizoplane while that of saprophytic fungi such as Aspergillus clavatus and T. viridae increase (Tables - 13,15,17 and 19). The <sup>4</sup>more increase in the frequency of A. clavatus and T. viridae in the rhizosphere and rhizoplane of JG-74 (known for antibiosis (Wakssman, 1952 and Gukulapalam and Nair, 1984)) might be one of the reason<sup>5</sup> for poor infection in the cultivar JG-74. Diathan Z-78 (200 ppm) has been found to influence the exudation of amino acids from the roots of Sorghum vulgare and Crotalaria jancea L., adversely resulting in the reduction in rhizosphere population (Balasubramanian and Rangaswami, 1973). It is likely that<sup>6</sup> similar mechanism might be operating here also. <sup>7</sup>In Antibiotic streptomycin, on the other hand, has been found to stimulate the rhizosphere fungal flora in the present studies. These results are in agreement with those of Gupta (1974), Chandra et al. (1982) and Bagyaraj and Rangaswami (1982). It is likely that antibiotic sprays reduce the bacterial counts in the rhizosphere, thus reducing the antibiosis or antagonistic effects on fungi which probably results in increase in soil rhizosphere fungi (Chandra et al., 1982). The possibility of the changes in the physiology and root exudation of the plants favouring growth of fungi in the



rhizosphere however, cannot be ruled out (Gupta, 1974 and Chandra et al., 1982).

Incorporation of different fertilizers in the form of urea, superphosphate, muriate of potash and oil cakes such as castor cake, neem cake, mustard cake results in an increase in the rhizosphere population of fungi in uninoculated plants of both the cultivars JG-62 and JG-74 while in inoculated ones, there has been a decrease in the rhizosphere activity except with urea and superphosphate. However, incorporation of mahua cake and different fungicides brings about a decrease in rhizosphere population of fungi in uninoculated and inoculated plants, of both the cultivars (Table - 20). These results are in agreement with those of Mosolove et al. (1959), Mishra (1972), Jarden et al. (1972), Jalaluddin (1975), Donech et al. (1983) and Tostensson and Wessen (1984). It is interesting to note that in the rhizosphere and rhizoplane of cultivar JG-62, F. oxysporum f.sp. ciceri was not detected in soil amended with mahua cake, neem cake, bavistin; while in JG-74 <sup>it was not detected after treatment</sup> with castor cake, mahua cake, neem cake, bavistin and benlate. The rhizosphere of inoculated plants is pre-dominated by parasitic/pathogenic fungi (Tables-21, 22, 23 and 24) which are adversely affected with muriate of potash, oil cakes and fungicides in the rhizosphere mycoflora (Jarden et al., 1972; Sinha, 1975; Haider et al., 1978; Singh et al., 1980; Upadhyay and Rai, 1981; Huang and Sun,

1982; Sulochna et al., 1983; Singh et al., 1985 and Sinha and Prasad, 1986).

It is likely that soil treatment with inorganic fertilizers may induce fresh root formation resulting in increase root exudation in the plants which provide<sup>s</sup> more substrate for rhizosphere and rhizoplane microorganisms. This might be one of the reasons for higher population<sup>s</sup> and frequency<sup>s</sup> of fungi. In inoculated plants, urea and superphosphate favour the activity of F. oxysporum f.sp. ciceri and other pathogenic forms (Tables-21,22,23 and 24). Similar results were obtained by Papendick and Cook (1974), Gamel-El-Din et al. (1983) and Rai and Upadhyay (1983).

Oil cakes, the product left after extraction of oils, contain ~~(sufficiently)~~ higher<sup>s</sup> amount<sup>s</sup> of lignin, cellulose and other carbohydrates, certain nitrogenous materials etc. which are suitable for all types of colonizers. <sup>Thereby</sup> thus result<sup>ing</sup> in ~~the~~ increase<sup>s</sup> of fungal population<sup>s</sup> particularly saprobes. Soil amendment with different oil cakes probably <sup>ok</sup> acts on pathogenic forms in many ways. Its decomposition products may be deleterious to the pathogenic forms, or it improves soil structure resulting in better growth of plants with modified root exudation unfavourable for pathogenic forms, or the degradation products which when exuded <sup>out</sup> ~~on~~ reduce the activity of parasitic forms (Huber and Walson, 1970; Kirmani, 1977; Upadhyay and Rai, 1981; Son et al., 1985

Nishat Khalis and Manoharachary, 1985 and Sinha and Prasad 1986. Lastly there may be <sup>a</sup>phenomenal increase in antagonistic forms of fungi such as T. viridae (Evans, 1955; and Saksena, 1960). In the present studies also there has been an increase in the frequency of T. viridae in oil cake amended soil which might also be contributing towards poor development of wilt. There has also been an increase in the frequency of <sup>species of</sup>Aspergilli and Penicilli in the rhizosphere<sup>s</sup> of oil cake amended soil. <sup>the</sup>Inhibitory effect of culture filtrate of A. niger is already known against F. oxysporum f.sp. ciceri (Mathur and Chauhan, 1972), Sinha and Prasad (1986) reported that castor cake and soybean leaves were more effective in reducing chickpea wilt disease. Similar results on the reduction of parasitic forms due to soil amendment with Azadiracta indica have been reported by Singh et al. (1980).

<sup>the</sup>Inhibitory effect on soil microorganisms as a result of soil treatment with pesticides <sup>have</sup> been reported by Bollen et al. (1954), Bollen (1961), Lebed (1964), Tu (1972 and 1973), Singh and Prasad (1973), Midhu and Nandwana (1974), Bertoldi et al. (1977 and 1978), Jain and Sehgal (1980), Sinha et al. (1980) and Bollen et al. (1983). The decrease in the activity of parasitic fungi with the amendment of fungicides might be due to <sup>the</sup>decrease in competitive saprophytic colonization of parasitic forms or increase<sup>s</sup> in

saprophytic forms (Vaartaja and Agnihotri, 1970 and Rai and Upadhyay, 1983). Moreover, changes in soil microbial enzyme activity cannot be ruled out (Munchi et al., 1988).

It, therefore, appears that the rhizosphere and rhizoplane mycoflora of chickpea is influenced by several factors. The rhizosphere and rhizoplane mycoflora of chickpea plants inoculated with Fusarium oxysporum f.sp. ciceri <sup>is</sup> different from those uninoculated ones, ~~whereas~~ In the former, the pathogenic forms predominate while in the later saprophytic forms <sup>predominate</sup>. Highest activity of rhizosphere/rhizoplane mycoflora is observed at flowering and fruiting stage<sup>s</sup> and not at senescence. Different cultivars differ in the rhizosphere and rhizoplane mycoflora both qualitatively and quantitatively. Sprays with various fertilizers, growth promoting substances, pesticides and soil treatment with different fertilizers, oil cakes, <sup>and</sup> fungicides influenced the rhizosphere and rhizoplane mycoflora in many ways. The frequency of Trichoderma viridae has been considerably increased due to application of oil cakes while parasitic forms such as Fusarium oxysporum f.sp. ciceri is decreased. These changes in the mycoflora appear<sup>s</sup> to be influenced by the root exudates which are affected by various treatments. The present studies go a long way in exploring and envisaging different possibilities where the rhizosphere mycoflora could be modified by foliar applications and soil amendment for control of

This statement is not valid since only minor differences were shown (not enough to affect control over a season) and these were not shown to be statistically significant differences verified by repeating the experiments.

root diseases. Brian (1957) also mentioned that successful control of root diseases probably lies in the development of satisfactory methods for influencing the rhizosphere microflora.

## SUMMARY

Rhizosphere<sup>S</sup> of uninoculated chickpea plants and those inoculated with Fusarium oxysporum f.sp. ciceri harboured more fungal population than ~~non-rhizosphere~~<sup>soil</sup>. Higher fungal population<sup>S</sup> was<sup>was</sup> encountered in the rhizosphere<sup>S</sup> of inoculated plants than in uninoculated ones. The concentration of total free amino acids, phenols, O-dihydroxy phenols and sugars was also high in the root extract of inoculated chickpea plants. By and large, the number of fungi was more in inoculated plants than in uninoculated ones. In the rhizosphere and rhizoplane of inoculated plants, parasitic fungi, particularly F. oxysporum f.sp. ciceri, dominated while saprophytic forms<sup>V</sup> in uninoculated plants.  
*dominate!*

2. There <sup>have</sup> been both qualitative and quantitative differences in the rhizosphere and rhizoplane mycoflora of different cultivars irrespective of the fact of their being inoculated with F. oxysporum f.sp. ciceri. Cultivar H-208 harboured <sup>the</sup> highest number of fungi in the rhizosphere while <sup>had the highest numbers</sup> H-208 and JG-62<sup>1</sup> in the rhizoplane. The fungus population was also high in the cultivar H-208 and least in JG-74 in both inoculated and uninoculated plants. There <sup>have</sup> been differences with regard to concentration of total free amino

acids, phenols, O-dihydroxy phenols and sugars in the roots of different cultivars with higher concentration<sup>s</sup> in inoculated plants than in uninoculated ones. The frequency of F. oxysporum f.sp. ciceri was also high in the rhizosphere and rhizoplane of cultivar JG-62, which supported more abundant rhizosphere and rhizoplane mycoflora, and least in JG-74, being less supportive to the rhizosphere and rhizoplane mycoflora.

3. The population of fungi in the rhizosphere increased with increase in age of plant<sup>s</sup> upto flowering and fruiting stage<sup>s</sup> followed by decline. The frequency of F. oxysporum f.sp. ciceri also increased with increase in age of the plants and attained<sup>the</sup> highest value when plants were 75 days old. The concentration of total free amino acids, phenols, O-dihydroxy phenols and sugars was highest at the age of 75 to 90 days old plants. By and large, Aspergillus<sup>sp.</sup> constituted the dominant mycoflora throughout the growth period in uninoculated plants. On the other hand F. oxysporum f.sp. ciceri and other parasitic fungi dominated in inoculated plants.

4. Indole acetic acid, indole butyric acid, thio-indole butyric acid, gibberellic acid, urea and streptomycin when applied as foliar sprays, showed inhibitory effect on the rhizosphere and rhizoplane mycoflora of inoculated and uninoculated plants of both the cultivars JG-62 and JG-74;

however, sprays with maleic hydrazide, muriate of potash, bavistin, vitavax, brassicol, benlate, captan and wettable sulphur exhibited inhibitory effect. The frequency of F. oxysporum f.sp. ciceri increased in the rhizosphere and rhizoplane when sprayed with indole acetic acid, indole butyric acid, thio-indole butyric acid, gibberellic acid, urea and streptomycin and decreased with maleic hydrazide, muriate of potash and different fungicides. F. oxysporum f.sp. ciceri was not isolated from rhizosphere and rhizoplane of both the cultivars when plants were sprayed with bavistin in the rhizosphere and rhizoplane of JG-74 and rhizosphere of JG-62 when plants were sprayed with benlate.

5. Soil amendment with different fertilizers in the form of urea, superphosphate, muriate of potash and oil cakes such as castor cake, neem cake, <sup>and</sup> mustard cake resulted in an increase in the rhizosphere population of fungi in uninoculated plants of both JG-62 and JG-74, but in inoculated ones there was a decrease in the rhizosphere population in all of the treatment except with urea and superphosphate. The soil amendment with mahua cake and with different fungicides brought about a decrease in rhizosphere population of fungi both in inoculated and uninoculated plants of the two cultivars. There was an increase in the frequency of F. oxysporum f.sp. ciceri in soil amendments with urea and superphosphate, but decrease in the remaining <sup>treatments</sup> F. oxysporum



f.sp. ciceri was either not detected or its frequency was low in the rhizosphere and rhizoplane of the plants.

## REFERENCES

- <sup>+</sup>Absalyamova, R.A. (1963). Effect of plant growth of the complex of root microorganisms occurring as a result of application of organo-mineral mixtures. Agrobiologia, 1:77-81 (Abstr. in Soil Fertilizers, 26:252, 1963).
- <sup>+</sup>Abdel-Kader, M.I.; Abdel-Fathah, H.M. and Hemida, S.K. (1983). Antifungal effect of bavistin, cotoran and curacron when incorporated with agar or liquid medium. Egypt. Jour. Bot., 24(2):143-148.
- <sup>+</sup>Afifi, A.F. (1976). Volatile and gaseous exudates of either germinating seeds or roots of Okra plants in relation to Okra wilt disease caused by some Fusarium species. Zentralblatt für Bakteriologie, Parasitenkunde, Infektions Krankheiten und Hygiene, 2 (1976), 131(6):555-564.
- Agarwal, A.K. (1986). Screening of soil fungi for antagonistic activity against soil borne plant pathogens (Abst.), Proc. All India Bot. Conf., :25.
- Agnihotrudu, V. (1953). Soil conditions and root diseases VIII. Rhizosphere microflora of some of the important crop plants of South India. Proc. Indian Acad. Sci. Sect. B, 37(1):1-13.
- Agnihothrudu, V. (1957). Fungi isolated from rhizosphere, III. Jour. Indian Bot. Soc., 36(4):486-490.
- Agnihothrudu, V. (1958). Fungi isolated from rhizosphere, IV. Jour. Indian Bot. Soc., 37(3):422-431.
- ✓ Agnihothrudu, V. (1959). Fungi isolated from rhizosphere, V. Jour. Indian Bot. Soc., 37:422-431.
- Agnihothrudu, V. (1960). Fungi isolated from rhizosphere, VI. Jour. Madras Univ. Sect. B, 29(3):155-181.

- Agnihothrudu, V., Bhuvaneshwari Nissk and Suryanarayanan, S. (1955). Fungi isolated from rhizosphere, I. Proc. Indian Acad. Sci. Sect. B, 42(3):98-104.
- Agnihotri, V.P. (1964). Studies on Aspergilli, XIV, Effect of foliar spray of urea on the Aspergilli of the rhizosphere of Triticum vulgare L. Plant and Soil 20(3):364-370.
- Allen, B.K. and Allen, O.N. (1958). Biological aspects of symbiotic nitrogen fixation. Encycl. Plant Physiology, 8:48-118.
- Alconero, R. and Hagedorn, D.J. (1968). The persistance of dexton in soil and its effect on soil microflora. Phytopath., 58(1):34-40.
- Annapurna, Y. and Rao, P.R. (1983). Foliar spray of urea and rhizosphere microflora of corn (Zea mays L.). Phytopathologische Zeitschrift, 106(2):104-107.
- Anon (1968). Antagonists to Sphacelotheca reiliana Trudy. Vologoel Moloch. Inst., 56:149-155 (Abstr. in Referal Zh. Rasten, 10:817, 1968).
- Ansari, A.R. (1982). Studies on the rhizosphere and rhizoplane mycoflora of sorghum and borley. Ph.D. Thesis, Aligarh Muslim University, Aligarh, India.
- ✓ Ansari, A.R. and Prakash, D. (1985). Studies on rhizosphere and rhizoplane mycoflora of Hordeum vulgare in relation covered smut disease. Proc. 72nd. Indian Sci. Cong. Part III.
- <sup>+</sup>Arora, D.K. (1985). Chemotoxis of Actinoplanes missouriens zoospores to fungal conidia, chlamydospores and sclerotia. Jour. Gen. Microbiol., 132(6):1657-1663.
- <sup>+</sup>Arora, D.K. and Dwivedi, R.S. (1979). Rhizosphere fungi of lens esculenta moench antagonistic to Sclerotium rolfsii Sacc. Soil Bio. and Biochem., 11(6):563-566.
- ✓ Ashour, W.A., Aly, A.A., Elewa, I.S. and Dabash, T. (1980). Interaction of soil microorganisms and rhizosphere of different onion cultivars with Fusarium oxysporum f.sp. cipae. The cause of basal rot disease of onion. Agri. Res. Review, 58(2):129-143.

- Atkinson, R.G. and Robinson, J.B. (1955). The application of a nutritional grouping method to soil fungi. Can. Jour. Bot., 33:281-288.
- Attabhanyo, A. and Holcomb, G.E. (1976). Control of Fusarium wilt of mimosa with systemic fungicides. Plant Disease Rep., 60(1):56-59.
- <sup>+</sup>Avezdzhanova, G.P., Matveev, G.G. and Tribunskii, A.N. (1974). [Microflora and the causal agent of wilt]. Microflora i vozbuditel Vilta. Khalopkovovodstvo, 8:24-25.
- <sup>+</sup>Babushkina, I.N. (1973). [Soil microscopic fungi of the rhizosphere of healthy and wilt infected cotton Gossypium hirsutum L. variety 18-F]. Pochvennye mikroskopicheskie griby rixofery khtopchalnika Gossypium hirsutum L. (Sort 108-F). Zolorovogo i porazheunogo verti tsillezhym viltom. Mikologiya i Fitopathologiya 7(6):525-531.
- <sup>+</sup>Babushkina, I.N. (1974). (Soil microscopic antagonistic fungi of the pathogenus of plant Verticillium wilt). Pochvennye mikroskopicheskie gribyantagonisty vozluditele i Vertitsilleznago vilta rostenii. Mikologia i Fitopathologia, 8(1):26-31.
- Bagyaraj, J. and Rangaseami, G. (1966). On the variation in rhizosphere effects of some crop plants. Curr. Sci., 35(9):238-239.
- Bagyaraj, J. and Rangaswami, G. (1967). Effect of fertilizers on the microflora of soil and the rhizosphere of Ragi (Eleusine coracana Gaertn). Indian Jour. Microbiol., 7(1):29-30.
- Bagyaraj, J. and Rangaswami, G. (1982). Rhizosphere microflora of Eleusine coracana as influenced by foliar chemical spray in the presence and absence of pathogen, Helminthosporium. Indian Phytopath., 35(2):388-392.
- Bahadur, P. and Sinha, S. (1965). Rhizosphere mycoflora of cluster beans (Cyamopsis psoraloides D.C.) in relation to root exudates. Proc. Nat. Acad. Sci. India, Sect. B. 35(3):343-350.
- Baker, R. and Nash, S.M. (1965). Ecology of plant in soil. VI. Inoculum density of Fusarium solani f.sp. phaseoli in bean rhizosphere as affected by cellulose and supplemented nitrogen. Phytopath., 55(12):1381-1382.

- Balasubramanian, A. and Rangaswami, G. (1973). Influence of foliar application of chemicals on the root exudation and rhizosphere microflora of Sorghum valgare and Crotalaria juncea. Folia microbiologia, 18(6): 492-498.
- Balasubramanian, A., Bagyaraj, D.I. and Rangaswami, G. (1970). Studies on the influence of foliar application of chemicals on the microflora and certain enzyme activities in the rhizosphere of Eleusine coracana Gaertn. Plant and Soil, 32(1):108-206.
- Barton, R. (1957). Germination of Oospores of Pythium mamillatum in response to exudates from living seedlings. Nature (Lond.), 180:613.
- Bartaria, A.M., Tamar, K.S. and Joshi, L.S. (1982). Fungicidal control of Sclerotial wilt of pan (Piper betle L.). Pesticides, 16(11):7-10.
- <sup>+</sup>Bateman, G.L. (1985). The effects of distribution of two soil incorporated fungicides on control of take all (Gaeumannomyces graminis Var. tritici) in wheat. Zeitschrift fur pflanzenkrankheiten und pflanzenschutz, 92(2):194-203.
- Bazhina, E.V. (1967). (Influence of herbicides on the development of soil fungi). Viganie gerbitsidov na rasutie pochvennykh gribov. Referat Zh. Bio., 12 B:487.
- Beggle Ristaino, J.E. and Papavizas, G.C. (1985). Survival and proliferation of propagules of Trichoderma spp. and Gliocladium virens in soil and in plant rhizosphere. Phytopath., 75(6):729-732.
- Behera, N. and Mukerji, K.G. (1979). Vertical distribution of fungi in two different soils of Delhi. Curr. Sci., 48(21):954-955.
- Behera, N. and Patnaik, R.S. (1980). Rhizosphere microflora of intact and defoliated plants of Dolichos biflavus Acta Bot. Indica, 8(1):105-109.
- <sup>+</sup>Berezova, (1941). Microflora of the rhizosphere of Flax. Microbiol. U.S.S.R., 10:918.
- Bergeson, G.B., Van Gundy, S.D. and Thomason, I.J. (1970). Effect of Meloidogyne javanica on rhizosphere microflora and Fusarium wilt of tomato. Phytopath., 60: 1245-1249.

- <sup>+</sup>Berestetskii, O.A. (1974). [Phytotoxic properties of microscopic fungi isolated from the rhizosphere of old apple trees]. Fitotoksicheskie svoystva mikroskopicheskikh gribov, vydelennykh iz rhizosfery stargkh derev'ev yabloni. Mikologiya i Fitopatologiya, 8(1):69-72.
- Bertoldi, M.De., Giovanunetti, M., Griselli, M. and Rambelli, A. (1977). Effect of soil application of benomyl and captan on the growth of onions and the occurrence of endophytic mycorrhizas and rhizosphere microbes. Ann. Appl. Biol., 86(1):111-115.
- Bertoldi, M.D., Rambelli, A., Giovanunetti, M. and Grisselli, M. (1978). Effect of benomyl and captan on rhizosphere fungi and growth of Allium cepa. Soil. Biol. Biochem., 10(4):265-268.
- Bhuvaneshwari, M. and Subba Rao, N.S. (1957). Root exudates in relation to the rhizosphere effect. Proc. Indian Acad. Sci., Sec. B, 45:299-301.
- Biehn, W.L. Lue, J. and Williams, E.B. (1968). Accumulation of phenols in resistant plant, fungi interactions. Phytopath., 58:1255-1260.
- Bisset, J. and Parkinson, D. (1979). Fungal community structure in some alpine soils. Can. Jour. Bot., 57(15):1630-1641.
- Block, R.J. (1950). Estimation of amino acids and amides on paper chromatography. Anal. Chem., 22:1327-1332.
- Bollen, W.B. (1961). Interactions between pesticides and soil microorganisms. Ann. Rev. Microbiol., 15:69-92.
- Bollen, W.H., Morrison, H.H. and Crowell, H.H. (1954). Effect of field treatment of insecticides on number of bacteria, Streptomyces and Molds in the soil. Jour. Ecol. Ent., 47:302-306.
- Bollen, C.J., Hoeven, E.P., Van Der, Lomers, J.G. and Schoonen M.P.M. (1983). Effect of benomyl on soil fungi associated with rye. II. Effect on fungi of culm bases and roots. Neth. Jour. Plant Path., 89(112):55-66.

- Bowen, G.D. (1979). Integrated and experimental approaches to the study of growth of organisms around roots. Pages 209-227. in: Soilborne Plant Pathogens. Schippers, B. and Gams, W. eds. Academic Press London 686 pp.).
- Boynton, D. (1954). Nutrition by foliar application. Ann. Rev. Plant Physiol., 5:31-54.
- Bray, H.G. and Thrope, W.V. (1954). Analysis of phenolic compounds of interest in metabolism. Meth. Biochem. Anal., 1:27-52.
- ✓ Brian, P.W. (1957). Ecological significance of antibiotic substances. In Microbial Ecology, 7th Symp. Soc. Gen. Microbiol. Cambridge University, Press.
- Bretage, T.W. (1985). Fungi associated with root rots of annual Medicago spp. in Australia. Trans. Brit. Myco. Soc., 85(2):329-334.
- Bretage, T.W. and Kollmorgen, T.F. (1986). Effect of triflurodin, benomyl and metalaxyl on the incidence and severity of root disease in annual Medicago spp. and evaluation of cultivar for resistance to root rot. Aust. Jour. Exp. Agr., 26(1):65-70.
- Bristow, P.R., and Wyllie, T.D. (1984). Reaction of soybean cultivars to Macrophomina phaseolina as seedlings in the growth chamber and field. Trans. of Missouri Acad. Sci., 18:5-10.
- Bumbieris, M. (1969). Effect of soil amendements on number of soil microorganisms and on root rot Fusarium wilt complex of peas. Aust. Jour. Bio. Sci., 22(6):1329-1336.
- Burden, R.S., Rogers, P.M. and Waijen, R.L. (1974). Investigations on fungicides VI. Natural resistance of plant roots to fungal pathogens. Ann. Appl. Bio., 78(1):59-63.
- Buxton, E.W. (1957). Differential rhizosphere effects of three pea cultivars on physiological races of Fusarium oxysporum f. pisi. Trans. Brit. Myco. Soc., 40(3): 305-316.
- Brown, S.L. and Curl, E.A. (1980). Rhizosphere effect of a herbicide stressed plant Cassia obtusifolia on Fusarium oxysporum f.sp. vesinfectum. Phytopath., 70(6):565.

- Cahill, D., Weste, G. and Grant, B. (1985). Leakage from seedling roots inoculated with Phytophthora cinnomomi. Phytopathologische Zeitschrift, 114(4): 348-364.
- Camporota, P. (1985). [Antagonism in vitro of Trichoderma spp to Rhizoctonia solani Kuhn]. Antagonisme in vitro de Trichoderma spp. vis-a-vis de Rhizoctonia solani Kuhn. Agronomie, 5(7):613-620.
- Carley, H.E. (1983). RH-5781F: A versatile new agricultural fungicide. The American Phytopathological Society (Southern Division). Ann. Meet. Feb. 6-9, 1983.
- Catska, V., Macura, J. and Vagnerova, K. (1960). Rhizosphere microflora of wheat III. Fungal flora of wheat rhizosphere. Folia Microbiol., 5:320-330.
- Chaube, H.S. and Singh, R.S. (1969). Relative abundance of hyphal and conidial propagules of Fusarium in rhizosphere and non-rhizosphere soil. Indian Phytopath., 22(2):273-275.
- Chauhan, S.K. (1963). Incidence of Fusarium wilt of Gram (Gier arietinum L.) in relation to soil moisture. Agr Univ. Jour. Res. (Sci.), 12(1):271-274.
- Chauhan, S.K. (1963). Incidence of Fusarium wilt of gram (Cicer arietinum L.) in oilcakes amended soil. Agra Univ. Jour. Res. (Sci.), 12(3):143-146.
- Chandra, S., Raizada, M. and Khanna, K.K. (1982). Changes in the rhizosphere microflora of tomato by foliar application of streptomycine. Indian Phytopath., 35(2):226-231.
- Chattopadhyay, S.B. and Mukhopadhyay, N. (1967). Investigation on disease complex of Rice in West Bengal (India). Int. Rice Res. Newsletter, 16(1):31-40.
- Chattopadhyay, A.K. (1984). Studies on the control of stem rot and root rot of Jute. Pesticides, 18(5): 25-27.
- Chao, W.L., Nelson, E.B., Harman, G.E. and Hoch, H.C. (1986). Colonization of the rhizosphere by biological control agents applied to seeds. Phytopath., 76(1): 60-65.



- Chawdhary, A.K. Mukhopadhyay, A.K. and Mukophadhyay, S. (1972). Influence of certain herbicides on fungal population in soil. Indian Phytopath., 25(2):188-194.
- Chester, C.G.C. and Parkinson, D. (1959). On the distribution of fungi in the rhizosphere of oats. Plant and Soil, 11(2):145-156.
- <sup>+</sup>Choroszewski, P.P. (1985). [Soil fungi and their pathogenicity to potato tubers] Grzyby Wystepujace W glebie i ich patogenicznosc W stosunku do bulwy ziemniaka. In Ziemniak. Bonin, Poland; Instytut Ziemniaka:117-134.
- Chrzanowski, J. (1976). [Mycoflora of the rhizosphere of wheat from field and in vitro investigations 1970-74]. Micoflora ryzosfery pszenicy Z badan polowych i in Vitro 1970-1974). Hodowla Roslin, Aklimatyzacja i Nasieunictwo, 20(1):19-45.
- Clark, F.E. (1940). Notes on types of bacteria associated with plant roots. Trans. Kansas Acad. Sci., 43:73-84.
- Clark, F.E. (1948). Rhizosphere microflora as affected by soil moisture changes. Soil Sci. Soc. Am. Proc., 12:239-242.
- Clark, F.E. (1949). Soil microorganisms and plant roots. Advances Agron., 1:24-88.
- Clark, F.E. and Thom, C.C. (1939). Effect of organic amendments upon the microflora of rhizosphere of cotton and wheat. Trans. 3rd Comm. Inst., Soc. Soil. Sci., A :94-100.
- Cochrane, W.V. (1958). Nitrogen Nutrition and Metabolism in Physiology of Fungi:241-299. Toppan Printing Company Ltd. Japan.
- Coley-Smith, J.R. and Hickman, C.J. (1957). Stimulation of Sclerotium germination in Sclerotium cepivorum Berk. Nature (London), 180:445.
- <sup>+</sup>Conci, V.C., Vasquez, A. and None, S.F. (1985). [Study of the form of transmission of Fusarium oxysporum f.sp. ciceri on chickpea (Cicer arietinum L.)]. Estudios de la forma de transmission de Fusarium oxysporum f.sp. ciceri engarbanzo (Cicer arietinum L.). Fitopatologia, 20(2):57-64.
- Cook, R.J. and Schroth, M.N. (1965). Carbon and nitrogen compounds and germination of chlamydospores of F. solani f.sp. phaseoli. Phytopath., 55:254-256.

- Collins, R.P. and Scheffer, R.P. (1958). Respiratory responses and systemic effects in Fusarium infected tomato plants. Phytopath., 48:349-355.
- Dath, A.P. (1982). Effect of soil amendment with some green manures on the survival of sclerotia of Corticium sasakii. Indian Jour. Phytopath., 35(3): 523-525.
- Davey, C.B. and Papavizas, G.C. (1960). Effect of decomposing organic soil amendments and nitrogen on fungi in soil and bean rhizosphere. Trans. Int. Cong. Soil Sci. 7th Congr. (Madison, Wist). Comm. III p. 551-557.
- David, M.B. and Baker, R. (1970). Rhizosphere competition in model soil system. Phytopath., 60:1058-1061.
- <sup>+</sup>Dayal, R. and Srivastava, L.S. (1975): Studies on the rhizosphere mycoflora of Abelmoscus esculentus Monch, I. Influence of varieties and age of the plant. Sydowia, 27(116):96-111.
- Davet, P., Ravise, A. and Baroudy, C. (1980). [The fungal microflora of bean (Phascolus vulgaris) roots in lebanon]. La microflora fungue des recines du horicotauliban. Ann.de Phytopathologie, 12(3):235-252.
- Deacon, J.W. and Mitchell, R.T. (1985). Toxicity of oat root extracts and saponins to zoospores of Pythium spp. and other fungi. Trans. Brit. Myco. Soc., 84(3): 479-487.
- Deb, P.R. and Dutta, B.K. (1964). Activity of thiolutin against certain soil borne plant pathogens. Curr. Sci., 53(12):659-660.
- Den Ouden, H. (1958). A new method for culturing plants enabling observation of nematodes on growing roots. Nath. Jour. Plant. Path., 64:269-272.
- Dick, M.W. and Ali-Shataych, M.S. (1986). Distribution and frequency of Pythium species in parkland and farmland soils. Trans. Brit. Mycol. Soc., 86(1):49-62.
- Dickinson, C.H. and Pugh, G.J.F. (1965a). The mycoflora associated with Helimione partulacoides I. The establishment of the root surface flora of mature plants. Trans. Brit. Mycol. Soc., 48:381-390.

- Dickinson, C.H. and Pugh, G.J.F. (1965b). The microflora associated with Helimione portulacoides II. Root surface fungi on the nature and excised plants. Trans. Brit. Mycol. Soc., 48:595-602.
- Doneche, B., Seguin, G. and Ribeau-Gayon, P. (1983). Mancozeb effect on soil microorganisms and its degradation in soil. Soil Sci., 135(6):361-366.
- Downes, M.J. (1972). Observations on some Fusarium populations in Co-Donegal oat soils. Scientific Proc. Royal Dublin Soc., B, 3(10):127-135.
- Dubois, M., Gilles, K., Hemilton, J.K., Robers, P.A. and Smith, F. (1956). Colorimeter method for determination of sugars and related substances. Anal. Chem. 28:350-356.
- Dutta, B.K. (1979). Studies on some fungi isolated from the rhizosphere of tomato plants and consequent prospect for the control of Verticillium Wilt. Proc. Indian Sci. Cong. 66th Session, III, Sect. 4, p.15.
- Dwivedi, R.S. and Mishra, R.C. (1982). Cladosporium cladosporioides as hyperparasite on hyphae of Rhizopus oryzae. Indian Phytopath., 35(4):704-705.
- Dwivedi, R.S. and Pathak, S.P. (1981). Effect of pesticides on non-target rhizosphere mycoflora of tomato. Proc. 68th Indian Sci. Congr. Part III. Section II, p. 23.
- Dwivedi, R.S. and Singh, B.P. (1971). Fungi in the root region of soybean (Glycine soja Linn.) I. Effect of foliar spray of gibberellic acid, maleic hydrazide and urea on the rhizosphere mycoflora. Trop. Ecol., 12(2):257-263.
- Elad, Y. Chet, I., Boyle, P. and Honis, Y. (1983). Parasitism of Trichoderma spp. of Rhizoctonia solani and Sclerotium rolfsii-Scanning electron microscopy and Fluorescence microscopy. Phytopath., 78(1):85.
- El-Hamalawi, Z.A. and Erwin, D.C. (1986). Components in alfalfa root extract and root exudate that increase Oospore germination of Phytophthora megasperma f.sp. medicaginis. Phytopath., 76(5):508-513.
- El-Hissy, F.T., Abdel Hafez, S.I. and Abdel Kader, N.I.A. (1983). Rhizosphere fungi of five plants in Egypt. Naturalis Monspeliensis, 34:11.

- Ellingboe, A.H. (1983). Genetical aspects of interaction between plant hosts and their soil borne pathogens. Phytopath., 73(6):941-944.
- Endo, S., Shinobara, N., Nakata, H., Kawata, N. and Yasuda, S. (1976). Studies on the soil borne pathogens parasitic to the same host. III. The reciprocal antagonistic actions of Corticium rolfsii (Sacc.) Curzi and Fusarium oxysporum schiechtendabl f. lycopersici (Sacc.) Snyder and Hansen and these effects on the occurrence of the diseases of tomato plants. Bull. of the Coll. of Agricond Vetekn Ned. Nihon Univ., 33:131-152.
- <sup>+</sup>Ernekova, B.D., Abirova, A.A. (1986). [Antagonistic properties of soil fungi in the Alma Ata district with respect to the causal agent of root rot] Izvestiya Akademii Nauk Kazakhskoi SSR, Biologicheskaya, No.2: 88-89.
- Evans, E. (1975). Survival and recolonization by fungi in soil treated with formation or carbon disulphide. Trans. Brit. Mycol. Soc., 38:335-346.
- Fenwick, L. (1973). Studies on rhizosphere microflora of onion plants in relation to temperature changes. Soil Bio. and Biochem., 5(3):315.
- Fries, N. and Forsman (1951). Quantitative determination of certain nucleic acid derivatives in pea root exudate. Physiol. Plantarum, 4:410-420.
- Funik - Jensen, D., and Hockenhull, J. (1984). Root exudation, rhizosphere microorganisms and disease control. Vaxtskyddsnotiser, 48(3-4):49-54.
- Furgal-Wegrzycka, H. (1984). [Research on the resistance of pea and field pea lines to pathogenic fungi with special reference to Fusarium and Ascochyta species. Part II. Research on the effect of water extracts of resistant and susceptible pea and field pea lines on the growth and development of pathogenic fungi]. Badania nad odpornoscia linii hodowlanych grochu i peluszek na porazenie przez chorobotworcze grzyby Ze 'szczegilnym uwzglednieniem gatunkow z-vodzajaiv Fusarium i Ascochyta. Czesi II. Badani nad wplywem wycia zeszyty Naukowe Akademi Rolnictwo, 39:137-153.
- Furuya, H.V.T. (1981). The significance of soil microorganisms on the inhibition of the macroconidial germination of Fusarium solani f.sp. phaseoli in a soil suppressive to common bean root rot. Ann. Phytopath. Soc. Japan, 47(1):42-49.

- Gadzhieva, M.A. (1959). The importance of organomineral mixtures and their component for the development of the root microflora of winter wheat and grass mixtures. Agrobiologiya, 4:581-589.
- Gagne, S., Autoun, H. and Richard, K. (1985). [Inhibition of plant pathogenic fungi by bacteria isolated from soil and the rhizosphere of leguminous plants]. Inhibition de champignons phytopathogènes par des bactéries isolées du sol et de la rhizosphère de légumineuses. Can. Jour. Microbiol., 31(9):856-860.
- Gamal-El-Din, H., Badar El-Din, S.M.S., Eld, E.T. and Makram, E.A. (1983). Quantitative changes in soil rhizosphere and phyllosphere microflora of plants as affected by nitrogen fertilization. Egypt. Jour. Microbiol., 18(1/2):1-7.
- Gangawane, L.V. (1985). Crop management and rhizosphere mycoflora, a review. Indian Bot. Rep., 4(2):160-168.
- Gangawane, L.V. and Deshpande, K.B. (1975). Agronomic treatments and changes in the rhizosphere mycoflora of groundnut IV. Effect of crop rotation. Proc. Indian Sci. Acad. B., 41(5):475-497.
- Gangawane, L.V. and Deshpande, K.B. (1977). Seasonal variation in the rhizosphere mycoflora of groundnut. Jour. Indian Bot. Soc., 56(4):289-295.
- Garret, S.O. (1970). Pathogenic root infecting fungi. Indian Phytopath., 36(3):561.
- Gerhardson, B., Alstrom, S. and Ramert, B. (1985). Plant reactions to inoculation of roots with fungi and bacteria. Phytopathologische Zeitschrift, 114(2):108-117.
- Gilligan, C.A. (1979). Modeling rhizosphere infection. Phytopath., 69(8):782-784.
- <sup>†</sup>Girczak, N. and Gierczak, Cz. (1973). Influence of fungicides on quantitative and qualitative composition of microflora in the soil of forest nursery at Rokownia and intensity of soil respiration. Bulletin dela, Societe, des Amisdes Sciences destetters de poznon D., 14:139-150.
- Goel, S.K. and Mehrotra, R.S. (1974). Rhizosphere and rhizoplane studies of Abelmoscus esculentus in relation to root rot disease. Indian Jour. Myco. and Plant Path., 4(11):40-48.

- Goodman, R.N., Kiraly, Z. and Zaitlin, M. (1967). The Biochemistry and physiology of infectious plant diseases. New Jersey: D. Van Nostrand Company Inc. Princeton.
- Gokulapalan, C. and Nair, M.C. (1984). Antagonism of few fungi and bacteria against Rhizoctonia solani Kuhn. Indian Jour. Microbiol., 24(1):57-58.
- Goos, R.D. and Timonin, M.I. (1962). Fungi from the rhizosphere of banana in Honduras. Can. Jour. Bot., 40:1371-1377.
- Gujrati, S. (1965). Crop management and rhizosphere of cultivated legumes. Ph.D. Thesis, Banaras Hindu University, Varanasi, India.
- Gujrati, S. (1968). Ecology of soil fungi of cultivated fields. Proc. Nat. Acad. Sci. India Sec. B., 38(1): 66-80.
- ✓ Gujrati, S. (1969). Effect of plant age on rhizosphere of micro-fungi. Proc. Nat. Acad. Sci. India Sec. B., 38(2):90-112.
- Gujrati, S. (1969). Rhizosphere fungi of two cultivated legumes. Proc. Nat. Acad. Sci. India Sec. B., 38(3-4):275-280.
- Gunasekar, V. and Rao, C.G. (1982). Effect of foliar application of the fungicides on the microflora of rhizosphere and non-rhizosphere soil of groundnut. Indian Phytopath., 36(1):203.
- Gupta, P.C. (1971). Foliar spray of gibberellic acid and its influence on the rhizosphere and rhizoplane mycoflora. Plant and Soil, 34:233-236.
- Gupta, V.K. (1974). Effect of foliar application of subamycine on rhizosphere and rhizoplane mycoflora Indian Phytopath., 27:267-268.
- Gussin, E.J. and Lynch, J.M. (1983). Root residues: substrate used by Fusarium culmorum to infect wheat, barley and ryegrass. Jour. Gen. Microbiol., 129(2): 271-275.
- Haider, M.G., Singh, R.K., Prasad, H., Nath, R.P. and Sharma, R.N. (1978). Effect of some common fungicides on the incidence of pigeon pea wilt. Indian Phytopath., 31(4):511-512.
- Halleck, F.E. and Cochrane, V.W. (1950). The effect of fungistatic agents on the bacterial flora of the rhizosphere. Phytopath., 40:715-718.

- Harley, T.L. and Waid, J.S. (1955). A method of studying active mycelia on living roots and other surface in soil. Trans. Brit. Myco. Soc., 38:104-118.
- Harper, J.L. (1950). Studies in the resistance of certain varieties of banana to Panama disease. I. Internal factor for resistance and antibiotics. II. The rhizosphere. Plant and Soil, 2(4):374-394.
- Harrison, N.A. and Beckman, C.H. (1982). Time/space relationship of colonization and host responses in wilt resistance and wilt susceptible cotton (Gossypium) cultivars inoculated with Verticillium and Fusarium oxysporum f.sp. vasinfectum. Physio. Plant Path., 21(2):193-207.
- Haware, M.P. and Nene, Y.L. (1982). Symptomless carrier of the chickpea wilt Fusarium. Plant Disease, 66(3): 250-251.
- Haware, M.P., Nene, Y.L. and Rajeshwari, R. (1978). Eradication of Fusarium oxysporum f.sp. ciceri transmitted in chickpea seed. Phytopath., 68(9):1364-1367.
- <sup>+</sup>Herr, L.J. (1957). Soil microflora associated with continuous cropping of corn, oats and wheat. Ohio Jour. Sci., 57(4):203-211.
- <sup>+</sup>Hiltner, L. (1904). Uber neuere Erfahrungen und problem auf dein Gebiet der Bodenbakteriologie und unter besonderer Berucksichtigung der Grundung und Brache. Arb. dtsh. Landw., Ges., 98(4):59-78.
- Hong, C.Y. (1969). Studies on Fusarium disease of crops. 1. The relation between rhizosphere fungi and the occurrence of damping off of cucumber seedlings. Effect of bacteria and actinomycetes isolated from the rhizosphere of cucumber seedling upon the occurrence of clamping off. Ann. Phytopath. Soc. Japan, 55(4):308-318.
- <sup>+</sup>Hongyal-Balul, W. (1983). [Fungal diseases of robinia seeds and seedlings and possibilities of their control]. Akaemages csiracsemet ek gambas betegegei es z elleniik valo vedekezes lehetosegei. Erdeszeti kutatasbk, 74: 343-349.
- Hornby, D. and Brown, M.E. (1977). Nitrate and ammonium in the rhizosphere of wheat crops concurrent observations of take-all. Plant and Soil, 48(2):455-471.

- Hornby, D. and Ullstrup, A.J. (1967). Fungal populations associated with maize roots. Composition and comparison of mycofloras from genotypes differing in root rot resistance. Phytopath., 57(8):869-875.
- Horst, R.K. and Herr, L.J. (1962). Effect of foliar urea treatment on number of actinomycetes antagonistic to Fusarium roseum f. cerealis in the rhizosphere of corn seedlings. Phytopath., 52:423-427.
- Huang, J.W. and Sun, S.K. (1982). The effect of nitrogenous fertilizers on disease development of watermelon Fusarium wilt. Plant Protection Bull., Taiwan, 24(2): 101-110.
- Huber, D.M. (1979). The role of nutrients in resistance of plants to disease. In Handbook of Nutrition and Food Vol.II (Edited by Recheigel, M., CRC Press, Palm Beach, Florida, U.S.A.).
- Huber, D.M. and Watson, R.D. (1970). Effect of organic amendments on soil borne plant pathogens. Phytopath., 60:22-25.
- Hus, S.C. and Lockwood, J.L. (1984). Biological control of Phytophthora root rot of soybean by Hypochytrium catenoides in greenhouse tests. Phytopathologische Zeitschrift, 109(2):139-146.
- Hussey, R.S. and Rancadori, R.W. (1977). Vertical distribution of soil microorganisms following subsoiling in a cotton management system. Phytopath., 67(6):783-786.
- Ingham, E.R. (1985). Review of the effect of 12 selected biocides on target and non-target soil organisms. Crop. protection, 4(1):3-32.
- Isarlishvilli, S.Y. (1957). Study of the fungal flora in the rhizosphere of Grape vines. Trud. Inst. Zeshch. Rest. Acad. Sci., Georgia, 12:125-136.
- Ishizawa, S., Suzuki, T., Sato, O. and Toyoda, H. (1957). Studies on microbial population in the rhizosphere of higher plants with special reference to method of study. Soil Plant Food, 3(2):85-94.
- <sup>+</sup>Ito, I. and Ui, T. (1975). Behaviour of Fusarium solani f.sp. phaseoli in the rhizosphere and under ground tissues of susceptible and non-susceptible plants. Memoir of the Faculty of Agriculture, Hokkaido Univ., 9(2):187-192.

Soiling?  
see p. 141



- Ivarson, K.C. and Katznelson, N. (1960). Studies on the rhizosphere microflora of yellow birch seedlings. Plant and Soil, 12:30-40.
- Jain, R.K. and Sehgal, S.P. (1980). Effect of certain pesticides on soil microflora. Jour. Myco. Plant Path., 10(1):88-89.
- Jalali, B.L. and Suryanarayana, D. (1972). Biochemical nature of root exudates in relation to the pathogenesis in root rot of wheat. Indian Phytopath., 25:195-199.
- Jalaluddin, M. (1975). Study of microorganisms from root regions of rice. Plant and Soil, 43(2):337-345.
- Johnson, G. and Schall, L.R. (1954). Relation of chlorogenic acid to the scab resistance in potatoes. Science, 115:627-629.
- Johnson, L.F. (1957). Effect of antibiotics on the numbers of bacteria and fungi isolated from soil by dilution plate method. Phytopath., 47:630-631.
- Jordan, V.W.L. (1975). Control of Verticillium wilt transmission in strawberry runners with systemic benzimidazole type fungicides. Ann. Appl. Biol., 81(1):43-50.
- Jordan, V.W.L., Such, B. and Eddy, B.P. (1972). Influence of organic soil amendments on Verticillium dahliae and on the microbial composition of the strawberry rhizosphere. Ann. Appl. Biol., 70(2):139-148.
- Kageyama, K. and Ui, T. (1982). Survival structure of Pythium species in the soils of bean fields. Ann. Phytopath. Soc. Japan, 48(3):308-313.
- Kamal and Singh, C.S. (1969). Rhizosphere mycoflora of certain ornamental plants. Sci. Cult., 35(11):635-637.
- Kamal and Singh, N.P. (1974). On microfungi from root region of ten sugarcane varieties. Indian Phytopath., 27(3):347-354.
- Kannaiyan, S. and Prasad, N.N. (1981). Changes in the rhizosphere microflora of rice on infection with Rhizoctonia solani. Oryza, 18(4):246-247.
- Kao, C.W. and Ko, W.H. (1986). Suppression of Pythium splendens in a Hawaiian soil by calcium and microorganisms. Phytopath., 76(2):215-220.

- <sup>+</sup>Karimbaeva, L. Ya and Sizova, T.P. (1976). [Mycoflora of the rhizosphere of some trees]. Omilcoflora rhizofery nekotorykh drevesnykh porod. Vestnik Moskovskogo Universiteta 6, 31(5):24-29.
- <sup>+</sup>Karimbaeva, L. Ya and Sizova, T.P. (1977). [Effects of root exudates of pine and spruce seedlings on the formation of mycoflora in the rhizosphere of these conifer species]. Vliyanie Kornevyskh vydelenic prorostkov sonsy i eli na formirovanie mikoflory rizosfery itykh porod. Mikologiy i Fitopatologiya, 11(3):191-199.
- Katznelson, H. (1946). The rhizosphere effect of mangels on certain groups of soil microorganisms. Soil Sci., XII(5):343-354.
- Katznelson, H. (1961). Microorganisms in the rhizosphere. Vol. I, p.610-614. In Recent Advances in Botany. (9th Int. Cong. Bot., Montreal, 1959). University of Toronto Press, Toronto.
- Katznelson, H. (1965). Nature and importance of the rhizosphere In Ecology of Soil-Borne Plant Pathogens (Eds. Baker, K.P. and Snyder, W.C.). Univ. California Press, Berkeley.
- Katznelson, H. and Richardson, L.T. (1948). Rhizosphere studies and associated microbiological phenomena in relation to strawberry root rot. Sci. Agr., 28(7): 293-308.
- Katznelson, H., Lochhead, A.G. and Timonin, M.I. (1948). Soil microorganisms and the rhizosphere. Bot. Rev., 14(9):543-587.
- Katznelson, H., Rouatt, J.V. and Payne, T.M.B. (1954). Liberation of amino acids by plant roots in relation to desiccation. Nature (Lond.), 174:1110-1111.
- Khan, M.A.A. (1984). Studies on rhizosphere and rhizoplane mycoflora of pigeonpea. Ph.D. Thesis, Aligarh Muslim University, Aligarh, India.
- Khan, M.A.A. and Prakash, D. (1982). Rhizosphere and rhizoplane mycoflora of gram affected by plant growth. Indian Phytopath., 35(4):717-718.

- ✓ Khan, M.W., Khan, A.M. and Saxena, S.K. (1973). The fungi and nematodes associated with roots of cauliflower. Indian Phytopath., 26(2):303-309.
- Khanna, R.N. and Singh, R.S. (1975). Microbial populations of pigeon pea rhizosphere in amended soil. Indian Jour. Mycol. Plant Path., 5(2):131-138.
- Khare, M.N. (1968). The relationship of Phytophthora fragariae Hickman to fungal flora in the rhizosphere of strawberry plant. Diss. Absor., 28(12)B:4836.
- +Khasanov, O.Kh. (1967). [Distribution of Penicillium spp. in the rhizosphere of Hibiscus cannabinus in a meadow marsh soil of chirckik river valley]. Rasprostraneniye gribov roda Penicillium Link v. rizofere kenafa na lugovobolotnoi pochve doliny r. chirckik. Uzbek. biol. Zh., 11(1):12-14.
- +Kirilenko, T.S. (1971). [Species of fungi of the family Dematiaceae in the rhizosphere of barley and oat in Ukrainian SSR]. Vidy gribov sem. Dematiaceae V rizofere yachmenya i Ovsa v Ukrainikoi SSR. Mikologiya i Fitopatologiya, 5(1):9-15.
- +Kirilenko, T.S. (1973). [The effect of fungi isolated from the rhizosphere of barley and oats on the growth of seedlings of these plants]. Vliyanie gribov, izolirovannykh iz rizofery yachmenya i ovsa, norost prorostkov etikh rastenii. Mikologiya i Fitopatologiya, 7(1):3-7.
- Kirmani, M.R. (1977). Studies on the role of fungi from decomposed oil-cakes on root-knot development on egg plants, Solanum melongena. Ph.D. Thesis, Aligarh Muslim University, Aligarh, India.
- Kirmani, M.R. and Alam, M.M. (1975). Influence of different C/N ratios on the population of fungi in the rhizosphere of cabbage. Indian Phytopath., 28(3):372-375.
- ✓ Kishan, S., Narendra, S. and Misra, S.R. (1982). Rhizosphere microflora associated with sugarcane plants affected by red rot. Indian Phytopath., 35(2):310-313.
- Knudson, L. (1920). The secretion of invertase by plant roots. Ann. Jour. Bot., 7:371-379.
- +Kotasthane, S.R., Agarwal, P.S. and Joshi, L.K. (1979). Studies on wilt complex in Bengal gram (Cicer orietinum L.). JNKVV Res. Jour., 10(3):257-258.

- Krassilnikov, N.V., Kriss, A.E. and Litvinov, M.A. (1936). The effect of root system on the soil microflora. Microbiol., 5(2):270-286. (English Summary).
- Krassilnikov, N.A., Raybalkina, A., Gabrillian, M. and Eondratievs, T. (1933). On the microbiological characteristic of the soil beyond the Volga region. Tran. Kromispo. Irrigacii. Acad. Nauk. U.S.S.R., 5:1.
- Krivets, I.O. (1975). [Erwinia carotovora in lupin rhizosphere]. Erwinia carotovora u rizadosferi lyupinu. Mikrobiol. Zh., 37(2):161-165.
- Kulshrestha, D.D. (1969). Studies on some soil fungi associated with maize (Zea mays L.) rhizosphere. Ph.D. Thesis, Aligarh Muslim University, Aligarh, India.
- Kulshrestha, D.D., Ray Chaudhari, S.P. and Khan, A.M. (1977). Studies on some soil fungi associated with maize (Zea mays L.) rhizosphere. I. Three new ascomycetes. Acta Bot. Indica, 5(1):16-19.
- Kumar, A.T.B. and Chakravarti, B.P. (1970). Changes in rhizosphere microflora of maize seedlings by pre-treatment of roots with hormones and urea. Plant and Soil, 33(3):679-684.
- Kumar, C.P.C. and Balasubramanian, K.A. (1981). Phyllosphere and rhizosphere microflora of pearl millet with reference to downy mildew incited by Sclerospora graminicola (Sacc.) Schroet. Plant and Soil, 62(1):65-80.
- Kundaswamy, D. and Rangaswamy, G. (1967). Changes in the rhizosphere microflora of sorghum due to foliar nutrient spray. Indian Jour. Agr. Sci., 2:143-150.
- <sup>+</sup>Kutrzeba, M. (1983). [Soil mycoflora as a factor limiting the occurrence of pathogenic fungi on three varieties of cocksfoot (Dactylis glomerata L.). Mikflora gleby jako czynnik ograniczajacy wystepowanie g rzybow pathogenicznych dla trzech odmian kupkowkipospolitej (Dactylis glomerata L.). Acta Mycologica, 19(2):245-281.
- <sup>+</sup>Kutrzeba, K. (1984). [Soil mycoflora as a factor limiting the occurrence of fungi pathogenic to three varieties of Dactylis glomerata L.II]. Mikoflora gleby jako czynnik ograniczajacy wystepowanie grzybow patogenicznych dla trzech odmian Dactylis glomerata L.II. Acta Mycologica, 20(1):33-44.

- Lacy, M.L. and Horner, C.E. (1962). Reproduction of Verticillium in the rhizosphere and subsequent root infection of resistant and susceptible Mint species. Phytopath., 52:739.
- <sup>+</sup>Lebed, E.S. (1964). [The effect of some herbicides on soil mycoflora]. Vliyaniye nekotorykh gerbitsidov na mikofloru pochvy. Nauch. Dokl. Yyssh. Shk, Bil. Sci., 1964(4): 171-176.
- <sup>+</sup>Lee, W.H., Ogoshi, A. and Sugimoto, T. (1985). Studies on the seed bacterization of sugarbeets.I. Comparative studies on the rhizosphere microflora of sugarbeets grown in different soils. Memoirs of the faculty of Agric. Hokkaido Univ., 14(4):405.
- Leonian, L.H. and Lilly, V.G. (1940). Studies on the nutrition of fungi.IV. Factors influencing the growth of some thiamin requiring fungi. Am. Jour. Bot., 27:18-26.
- Lewis, J.A. (1979). Influence of soil texture on survival and saprophytic activity of Rhizoctonia solani in soils. Can. Jour. Microbiol., 25:1310-1314.
- Lilly, V.G. and Barnett, H.L. (1951). The essential nonmetallic elements other than carbon in Physiology of the Fungi. pp.87-115. Mc Graw-Hill Book Company Inc. New York.
- Lim, L.L. and Cole, A.L.J. (1984). Infection and colonization of white clover seedling roots by fungi. Newzealand Jour. of Agric. Res., 27(3):459-465.
- Lochhead, A.G. (1940). Qualitative studies of soil microorganisms III. Influence of plant growth on the character of bacterial flora. Can. Jour. Res. Sect. C., 18(2): 42-53.
- Lochhead, A.G. (1952). Soil Microbiology, Ann. Rev. Microbiol., 6:185-206.
- Lochhead, A.G. (1959). Rhizosphere microorganisms in relation to root disease fungi. In Plant Pathology Problems and Progress (1908-1958). (Eds. Holton et al., Univ. Wisconsin Press, Madison, p. 327-338.
- Lockwood, J.L. (1985). Soil borne plant pathogens: concepts and connections. Phytopath., 76(1):20-27.
- Loper, J.E., Suslow, T.V. and Schroth, M.N. (1984). Lognormal distribution of bacterial population in the rhizosphere. Phytopath., 74:1454-1460.

- Louvet, J. (1972). Significance of quantitative and qualitative studies of phytopathogenic soil fungi. EPPQ Bulletin, 7:5-15.
- Luke, P. and Devi, S.S. (1975). Rhizosphere mycoflora of Pisum sativum L. New Botanist, 2(3/4):128-131.
- Luke, P. and Vani, M.A.M. (1972). Rhizosphere and rhizoplane mycoflora of Nicotiana tabacum L. Mysore Jour. Agric. Sci., 6(2):204-206.
- Lumsden, R.D., Lewis, T.A.<sup>L</sup> and Millner, P.D. (1983). Effect of composted sewage sludge on several soil borne pathogens and diseases. Phytopath., 73(11):1543.
- Lundegardh, H. and Stenlid, G. (1944). On the exudation of nucleotides and flavonones from living roots. Arch. Bot., 31A:1-27.
- <sup>+</sup>Lyon, T.L. and Wilson, J.K. (1921). Liberation of organic matter by roots of growing plants. New York Agric. Exp. Sta. (Geneva) Mem., 40:1-44.
- Malajeczuk, N., Boughton, T.J., Campbell, N.A. and Litchfield R.T. (1983). Interaction between the fungicide metalaxyl and soil microorganisms on survival of Phytophthora cinnamomi. Ann. Appl. Bio., 103(1):57-61.
- <sup>+</sup>Maliszewska, W. and Moreau, R. (1959). The rhizosphere of white spruce. Compt. Rend., 249:303-305.
- Mall, S. (1973). Studies on microbial antagonism in rhizosphere soils of potato varieties differentially susceptible to black-scurf and wilt. Plant and Soil, 39(3): 547-554.
- Manoharachary, C. and Bilolikar, V.G. (1979). A note on the distribution of mycoflora on some soils of border areas of Andhra Pradesh. Maharashtra Vidyan Mandir Patrika, 14(2):81-82.
- Manoharacharya, C., Venkatesharlu, K. and Rama Rao, P. (1977). Studies on mycoflora of rhizosphere and non-rhizosphere soils. Geobios., 4(2):67-68.
- <sup>+</sup>Maric, A., Balaz, F. and Jasnie, S. (1974). Fusarium wilt of watermelon (Fusarium oxysporum f. niveum) and possibilities of its control. Plant Protection (Zastita Bilja):15-27.

- Marimuthu, T. and Kandaswamy, T.K. (1980). Bacterial leaf blight of green gram (Vigna radiata Var. aureus) caused by Xanthomonas phaseoli and reaction of certain varieties to the disease. Madras Agric. Jour., 67:413.
- Martin, J.P. (1950). Use of acid, rose bengal and streptomycin in the plate method for estimating soil fungi. Soil Sci., 69:215-233.
- ✓ Mathur, S. and Chauhan, S.K. (1972). A comparative study of the rhizosphere of gram (Cicer arietinum L.) in relation to its wilt disease caused by different fungi. Jour. Indian Bot. Soc., 51(3/4):267-274.
- Mc Lean, R.S. and Cook, I.W.R. (1957). Practical field ecology. George Allen and Unwin Ltd. London.
- Meghraj, M., Prabhakara Rao, A. and Venkateswalu, K. (1987). Microbial population in soil from a subabul (Leucaena leucocephala) field. Acta Bot. Indica, 15:145-146.
- Mehrotra, R.S. and Claudius, G.R. (1973). Competitive saprophytic ability of Fusarium oxysporum f. lentis and Sclerotium rolfsii responsible for wilt and root rot disease of Lens culinaris at Sagar. Jour. Indian Bot. Soc., 52(3/4):191-198.
- Mew, T.W., Rosales, A.N. (1984). Relationship of soil microorganisms to rice sheath blight development in irrigated and dryland rice cultivars. Tech. Bull. ASPAC Food and Fert. Techno. Center, Taiwan No.79:11 (Abst. in Tropical Agric., 10:51252).
- Miller, R.E. and Bothroyd, C.W. (1962). Seasonal population of rhizosphere fungi associated with corn roots. Phytopath., 52:744.
- Mishra, R.R. (1972). Effect of certain chemical fertilizers on the rhizosphere mycoflora of Oryzae sativa Linn. III. Phosphate fertilizers-Superphosphate. Mycopath. Mycol. Appl., 46(1):97-102.
- Mishra, R.R. and Kamal (1972). Rhizosphere fungal flora of certain Euphorbiaceous plants. Mycopath. Mycol. Appl., 46(1):73-79.
- Misra, R.R. and Gujrati, S. (1969). Effect of maleic hydrazide on growth performance of certain fungal species. Sci. Cult., 35(3):109-110.

- Midha, S.K. and Nandwana, R.P. (1974). Effect of certain nematicides on the population of soil microorganisms. I. Quantitative and qualitative changes in fungal flora. Indian Phytopath., 27(3):312-345.
- Mikzak, M., Piotrowski, J. and Segit, Z. (1985a). [Differences in the chemical composition of borne hop forms depending on the degree of susceptibility to Verticillium albo-atrum and Fusarium sambucinum I. Content of phenol compounds]. Różnice Wskładzie chemicznym kilku form chmielu, w zależności od stopnia podatności na Verticillium albo-atrum i Fusarium sambucinum Część I. Zawartość związków fenolowych. Biuletyn Instytutu Hodowli i Aklimatyzacji Roslin. No.158:101-107.
- Mikzak, M., Piotrowski, J. and Segit, Z. (1985b). [Differences in chemical composition of some hop forms depending on degree of susceptibility of Verticillium albo-atrum and Fusarium sambucinum II. Content of free amino acids and soluble sugars and peroxidase activity]. Różnice Wskładzie chemicznym kilku form chmielu w zależności od stopnia podatności na Verticillium albo-atrum i Fusarium sambucinum Część II. Zawartość wolnych aminokwasów i cukrów rozpuszczalnych oraz aktywność peroksydaz. Biuletyn Instytutu Hodowli i Aklimatyzacji Roslin. No.158:109-115.
- Mohamed, Z.K. (1985). Physiological and antagonistic activity of Streptomyces in rhizosphere of some plants. Egypt. Jour. Phytopath., 14( $\frac{1}{2}$ ):121-128.
- Moor, H. and Stein, W.H. (1954). Modified ninhydrin reagent for the spectrophotometric determination of amino acids. Jour. Biol. Chem., 24:904-913.
- Moskovets, V.S. (1957). Fungal flora of the rhizosphere of lucerne in South Ukrainian SSR. Information. II. The quantitative and qualitative composition of the fungal flora of the rhizosphere of lucerne. Jour. Microbiol. Kiev., 19(3):44-50.
- Moskovets, V.S. and Zhdanova, N.M. (1960). The qualitative and generic composition of the fungal flora of the rhizosphere of maize in some regions of the steppe and forest steppe of the Ukrainian SSR. Jour Microbiol Kiev., 22(3):21-26.
- Moubasher, A.H. and Abdel-Hafez, S.I.I. (1986). Effect of soil amendment with three organic substances on soil, rhizosphere and rhizoplane fungi and on the incidence of damping off diseases of cotton seedlings in Egypt. Naturalia Monspeliensia, No.5:91-108.



- Mujumdar, S.B. and Bhide, V.P. (1970). Effect of carbon dioxide concentration on fungi from rhizosphere and non-rhizosphere soil of sugarcane.II. Persistence of rhizosphere effect in sugarcane soil. Jour. Univ. Poona Sci. Tech. Sect. 38: 137-141; 143-145.
- Mukhopadhyay, D. and Nand, B. (1974). Rhizosphere mycoflora of healthy and infected jute plants. Curr. Sci., 43(17): 590-591.
- Mukerji, K.G. (1968). Observations on the mutual relationship among soil microorganisms. Jour. Gen. Appl. Microbiol., Tokyo, 14(3):243-250.
- Munshi, M. Raghu, K. and Shah, A.M. (1988). Fungicidal effect on various soil microbial enzyme activity. Int. Conf. Res. Plant Sci. and its Relevance to Future Sec. V.pp 76.
- Murthy, N.B.K. and Raghu, K. (1976). Effect of thiram on plant growth and rhizosphere microflora of barley and nodulation in cowpea. Plant and Soil, 44(2):491-493.
- Mustafa, T.P. and Chatopadhyay, S.B. (1981). Reaction of cabbage varieties of Fusarium oxysporum f. conglutinans in relation to root exudates. Indian Phytopath., 32(2): 255-258.
- <sup>+</sup>Namoradize, I.I. (1975). Effect of different temperatures on the growth and development of Fusarium species isolated from the roots of peach. Trudy Nauchno Issledovatel, Skogo Instituta Zashchily Rostenii Cruz. SSR., 27:153-157.
- Narendra Singh and Singh, R.S. (1985). Inhibition of Fusarium udum (pigeonpea with pathogen) by ether distillate of Margosa cake amended soil. Indian Jour. Myco. and Plant Path., 13(3):329-330.
- Narendra Singh, Singh, R.P. and Lal, S. (1985). Effect of soil amendment on sugarcane wilt and its pathogens. Acta Bot. Indica, 13(2):212-217.
- Natrajan, K. (1972). An evaluation of some technique for isolation of rhizosphere and soil mycoflora. Proc. Indian Acad. Sci., Sect. B, 26(1):31-38.
- Neal, J.L., Atkinson, T.G. and Larson, R.I. (1970). Changes in the rhizosphere microflora of spring wheat induced by disomic substitution of a chromosome. Can. Jour. Microbiol., 16(3):153-158.

- Neate, S., Rovira, A. and Warcup, J.H. (1981). Minimum cultivation and root disease in wheat. Biannual Report of White Agri. Res. Inst. (1980-81).
- <sup>+</sup>Newman, E.I. (1978). Root microorganisms : Their significance in the ecosystem. Biological Reviews of the Cambridge Phylosophical Society, 53(4):511-554.
- Nishat Khalis and Manoharachary, C. (1985). Studies on the mycoflora changes in oil cakes amended and unamended soil. Indian Phytopath., 38(3):462-466.
- Obraztsova, A.A. (1935). The rhizosphere microorganisms of Batum red soils. Doklady Akad. Nauk. U.S.S.R., 94:70-71.
- Odunfa, V.S.A. and Oso, B.A. (1979). Fungal population in the rhizosphere and rhizoplane of cowpea. Trans. Brit. Mycol. Soc., 73(1):21-26.
- <sup>+</sup>Oku, H., Oki, K., Shiraish, T., Sato, K. and Ouchi, S. (1979). Effect of fungicides benomyl and thiram on soil microflora and some soil inhabitant fungi. Scientific Reports of the faculty of Agr. Okayama Univ., 54:1-8.
- Onofrash, L.F., Deshkova, A.D., Statrova, G.L., Bumistru, L.S., Lazu, M.N., Tinku, V.L. and Deviclovich, R.E. (1976). Effect of mineral fertilizers on the development of Verticillium disease of eggplant. Zefertiuny Zhurnal,: 68-74 (English summary).
- O'Rourke, C.J. and Millar, R.L. (1966). Root rot and root microflora of alfalfa as affected by potassium nutrition, frequency of cutting and leaf infection. Phytopath., 56(9):1040-1046.
- Padma, R. and Mukerji, K.G. (1972). Fungi in the root region of Rauwolfia serpentina and Rauwolfia canescens. Indian Phytopath., 25(2):104-107.
- Panwar, K.S., Panwar, S. and Bilgrami, K.S. (1967). Rhizosphere mycoflora in the sand-dunes of western Rajasthan. Proc. Nat. Acad. Sci. India B,
- Papavizas, G.C. and Davey, C.B. (1961). Extent and nature of the rhizosphere of lupinus. Plant and Soil, 14(3):215-236.
- Papavizas, G.C. and Kovacs, M.F. (1972). Stimulation of spore germination of Thielaviopsis basicola by fatty acid from rhizosphere soil. Phytopath., 62(7):688-694.

- Parker, M.A. (1985). Local population differentiation for compatibility in an annual legume and its host specific fungal pathogen. Evolution, 39(4):713-723.
- Parkinson, D. (1957). New methods for the qualitative and quantitative study of fungi in the rhizosphere. Symposium Methodes d' Etudes Microbiologiques du sol, June, 1957, Louvain, Belgium.
- ✓ Parkinson, D. (1967). Soil microorganisms and plant roots. In Soil Biology (by Buges, A. and Raw, F.) Academic Press, London and New York, pp. 449-478.
- Parkinson, D. and Clark, J.H. (1961). Fungi associated with the seedlings of Allium porrum L. Plant and Soil, 13: 384-390.
- Parkinson, D., Taylor, G.S. and Peason, R. (1963). Studies on fungi in the root region. Plant and Soil, 19:332-349.
- Parlik, M. (1950). [The microflora of the rhizosphere of Flax seedlings grown in sterile soil and in soil infested with Fusarium lini Bolley]. Mikroflora rhizosfery Khienich rostlinek Inu pestoranych v pude sterilni a infikowane' sporami Fusarium lini Bolley. Sporn. Csl. Akad. Zemed (Annl. Acad. lechecosl. Agric.), 23(3-4): 198-203. (Abst. in Biol. Abstr., 25(12):3352-3353.
- Patel, B., Fisher, J. and Wallace, H.R. (1981). Relation of soil borne organisms to disease in wheat. Biennial Report of Waite Agric. Res. Inst., p.198.
- Patel, B., Fisher, J. and Wallace, H.R. (1984). Relation of soil borne organisms to disease in wheat. Biennial Report of the Waite Agric. Res. Inst. p. 144.
- Patel, J.J. and Iyer, V.N. (1961). Microorganisms associated with the rhizosphere of cotton plant. Proc. Indian Acad. Sci., 44:1-10.
- Perotti, R. (1926). On the limits of biological inquiry in soil science. Proc. Int. Soc. Soil Sci., 2:146-161.
- Peterson, E.A. (1958). Observations on fungi associated with plant roots. Can. Jour. Microbiol., 4:257-265.
- Peterson, E.A. (1959). Seed borne fungi in relation to colonization of roots. Can. Jour. Microbiol., 5(6):579-582.

- Peterson, E.A. (1961). Observation on the influence of plant illumination on the fungal flora of roots. Can. Jour. Microbiol., 7:1-6.
- Peterson, E.A. and Timonin, M.I. (1977). Influence of amino acids and substances from flax roots on the soil borne plant pathogen Fusarium oxysporum f. lini. Can. Jour. Plant Sci., 57(2):407-411.
- Phillips, A.J.L. (1986). Factors affecting the parasitic activity of Gliocladium virens on Sclerotinia sclerotiorum and a note on its host range. Jour. Phytopath., 116(3): 212-220.
- Phipps, P.M. and Stipes, R.J. (1975). Control of Fusarium wilt of mimosa with benomyl and thiobendazole. Phytopath., 65(4):504-506.
- Pieczarka, D.J. and Lorbeer, J.W. (1975). Microorganisms associated with bottom rot of lettuce grown on organic soil in New York State. Phytopath., 65:16-21.
- Pineda, J.B. and Diaz Palanca (1981). [Biological control of Sclerotium rolfsii Sacc. on Phaseolus vulgaris using Penicillium notatum Westt.]. Control biologica de Sclerotium rolfsii Sacc. en Phaseolus vulgaris mediante La Utilization de Penicillium notatum West. Agromonia Tropical, 31(1/6):265-281.
- Prakash, D. (1967). Studies on population changes and ecology of fungi around the roots of sugarcane. Ph.D. Thesis, Aligarh Muslim University, Aligarh, India.
- Prakash, D., Ansari, A.R., Ashraf, S. and Khan, A.M. (1979). Studies on rhizosphere mycoflora of certain vegetables. Proc. 5th. Ann. Conf. Soc. Adv. Bot., p.25.
- ✓ Rai, B. and Upadhyay, R.S. (1980). The population dynamics of Fusarium udum Butler in soil in relation to root region microflora of healthy and wilted plants of pigeonpea. Proc. Nat. Acad. Sci. India Sect. Biol. Sci. p.68 (Golden Jubille Session).
- Rai, B. and Upadhyay, R.S. (1983). Competitive saprophytic colonization of pigeonpea substrate by Fusarium udum in relation to environmental factors, chemical treatments and microbial antagonism. Soil Biol. Biochem., 15(2): 187-191.

- <sup>†</sup>Raicu, C. and Stan, G. (1975). Soil fungi pathogenic to tomato. Interrelations with other microorganisms of the soil. Probleme de Protectia Plantelor., 3(3): 229-248.
- Ramachandra Reddy, T.K. (1959). Foliar spray of urea and rhizosphere microflora of Rice (Oryzae sativa L.). Phytopath. 4., 36(3):286-289.
- Ramachandra-Reddy, T.K. (1968). Plant treatment in relation to rhizosphere effect.I. Pretreatment of root of rice seedlings to certain chemicals and antibiotics on the rhizosphere microflora. Plant and Soil, 28:242-356.
- Rangaswami, G. and Vasantharajan, V.N. (1961). Studies on the rhizosphere mycoflora of citrus plants. as influenced by streptomycin spray. Curr. Sci., 30:25-26.
- ✓ Rangaswami, G. and Vasantharajan, V.N. (1962). Studies on the rhizosphere microflora of citrus trees.I. Quantitative incidence of microorganisms in relation to root and shoot growth.III. Fungal and actinomycete flora of rhizosphere. Can. Jour. Microbiol., 8:473-477.
- Ranga Rao, V. (1971). Studies on fungi in the root zone of cultivated plants (Part I) and cytology of ascus in chaetomieaceae (Part II). Ph.D. Thesis, University of Delhi, Delhi, India.
- Ranga Rao, V., Jayakar, M., Sharma, K.P. and Mukerji, K.G. (1972). Effect of foliar spray of morphactin on fungi in the root zone of Capsicum annum. Plant and Soil, 37:179-182.
- Rao, A.S. (1962). Fungal population in the rhizosphere of peanut (Arachis hypogaea L.). Plant and Soil, 17:260-266.
- Rao, A.V. and Sharma, R.K. (1980). Influence of chemicals on the microflora and enzyme activities in the cauliflower rhizosphere. Acta Bot. Indica, 6:71-74.
- Rao, R. and Raja, V. (1978). Effect of foliar application of urea and tetracycline on the rhizosphere microflora of wheat infected with Sclerotium rolfsii. Symp. Plant Dis. Prob., Jaipur, p.33-34.
- ⌣ <sup>†</sup>Rataj-Guranowsky, M. (1981). Communities of fungi from the soil, rhizosphere and rhizoplane surrounding roots of two varieties of yellow lupine (Lupinus luteus L.) susceptible and resistant to Fusarium wilt disease. Genetica Polonica, 22(1):9-23.

- Reddy, M.V. and Singh, K.B. (1981). Foliar application of Brovo 500 for Ascochyta blight control. Int. Chickpea Newsletter, No.8:25-26.
- ✓ Reddy, T.K.R. (1959). Rhizosphere microflora of pteridophytes. Curr. Sci., 28(3):113-114.
- Reis, E.M. (1982). [Effect of soil treatment with fungicides on seedling resistance, growth, common root rot and yield of wheat (Triticum aestivum)]. Efeito de tratamento de solo com fungicidas sobre a resistencia de plantulas, crescimento, podridao cumundadas raizes e rendimento de trigo (Triticum aestivum). Summa. Phytopathologica, 8(1/2):107-115.
- Renolds, R.S. (1926). Nutritional studies on Fusarium lini. Plant Physiol., 1:151.
- +Riker, A.J. and Riker, A.S. (1936). Introduction to research on plant disease. Plantographed by Johns. Swiftco Inc. Loutism Chicago, New York.
- Riviera, J. (1959). Contribution to the study of wheat rhizosphere. An. Agron., 45:293-337.
- Robinson, B.M. (1970). Micro-fungi of sugarcane roots and soil in Jamaica. Trop. Agric. Trin., 47(1):23-29.
- Rouatt, J.W. (1959). Initiation of rhizosphere effect. Can. J. Microbiol., 5:67-71.
- Rouatt, J.W. and Katznelson, H. (1960). Influence of light on bacterial flora of roots. Nature (Lond.), 186:659-660.
- ✓ Rouatt, J.W. and Katznelson, H. (1961). A study of the bacteria on the root surface and in the rhizosphere soil of crop plants. Jour. Appl. Bacteriol., 24:164-171.
- Rouatt, J.W., Katznelson, H. and Henderson, V.E. (1963). Micro-organisms in the root zone in relation to temperature. Can. Jour. Microbiol., 9:227-236.
- ✓ Rovira, A.D. (1956a). Plant root excretions in relation to the rhizosphere effect.I. The nature of root exudate from oats and peas. Plant and Soil, 7:178-194.
- ✓ Rovira, A.D. (1956b).III. The effect of root exudate on number and activity of microorganisms in soil. Plant and Soil, 7:209-217.

- ✓ Rovira, A.D. (1959). Root excretion in relation to the rhizosphere effect IV. Influence of plant species, age of plant, light, temperature and calcium nutrition on exudation. Plant and Soil, 11:53-64.
- ✓ Rovira, A.D. (1962). Plant root exudates in relation to the rhizosphere microflora. Soil Fertilizers, 25:167-172.
- ✓ Rovira, A.D. (1965). Interaction between plant roots and soil microorganisms. Ann. Rev. Microbiol., 19:241-266.
- ✓ Rovira, A.D. (1969). Plant root exudates. Bot. Rev., 35: 35-37.
- ✓ Rovira, A.D. and Harris, T.R. (1961). Plant root excretions in relation to rhizosphere effect.V. The exudation of B group vitamins. Plant and Soil, 14:119-214.
- Roy, R.Y. and Dwivedi, R.S. (1967). Rhizosphere studies of cultivated legumes. Proc. Indian Sci. Cong., p.294.
- Saksena, S.B. (1960). Effect of carbon disulphide fumigation of Trichoderma viridae and other soil fungi. Trans. Brit. Mycol. Soc., 43:111-116.
- Sakthivel, N., Sivamani, E., Unnamalal, N. and Gnanamanikam, S.S. (1986). Plant growth promoting rhizobacteria in inhancing plant growth and suppressing plant pathogens. Curr. Sci., 55(1):22-25.
- Salt, G.A. (1982). Factors affecting resistance to rot and wilt disease. In Fababean improvement. The Hague Netherlands, Mortinus, Nijhoff:259-220.
- <sup>+</sup> Samilinko, V.I. and Govozdyak, R.I. (1972). [Distribution of bacteriophages of Pseudomonas spp. causing legume diseases in the rhizosphere of crop plants]. Poshirnnya bakteriofagiv zoudnikiv bakterioziv bobovikh 4 rodu Pseudomonas V rhizofers sil's kogospodors'kikh roslin. Mikrobiologichnil Zhurnal, 34(4):518-520.
- Samtsevich, S.A. and Borisova, V.N. (1961). Effect of fertilizers on the root microflora of winter wheat. Microbiol., (USSR):30:1033-1041 (English Translation).
- ✓ Satyaprasad, K. (1982). Rhizosphere and rhizoplane mycoflora of wilt resistant and susceptible chickpea. Indian Phytopath., 35(1):153-155.

- Satyaprasad, K. and Rama Rao, P. (1982). Soil fungistasis against Fusarium oxysporum f.sp. ciceri causing wilt of chickpea. Proc. Indian Nat. Acad. Sci. B., 48(5): 652-658.
- ✓ Satyaprasad, K. and Rma Rao, P. (1983). Effect of chickpea root exudates on Fusarium oxysporum f.sp. ciceri. Indian Phytopath., 36(1):77-81.
- Schench, S. and Stotzky, G. (1975). Effect on microorganisms of volatile compounds released from germinating seeds. Can. Jour. Microbiol., 21(10):1622-1634.
- Scroth, M.N. and Hildebrand, D.C. (1964). Influence of plant exudates on root infecting fungi. Ann. Rev. Phytopath., 2:101-132.
- Scroth, M.N. and Snyder, W.C. (1961). Effect of host exudates on chlamydospores germination of bean root fungus, F. solani f.sp. phaseoli. Phytopath., 51:389-393.
- Sembdner, G., Gross, D., Liebisch, H.W. and Schneider, G. (1980). Biosynthesis and Metabolism of plant hormones in Encyl. Plant Physiol. Vol.9 (Edited by Mac Millan, J. New York, pp.281-390).
- Sharma, B.L. and Gupta, R.N. (1983). Survival of Fusarium oxysporum f.sp. ciceri causing wilt of Bengal gram during summer months. Indian Phytopath., 36(3):561-562.
- Sharma, K.P., Batta, S.K. and Kanwar, R.S. (1982). Phenolic contents in red rot resistant and susceptible varieties of sugarcane. Jour. of Res., Punjab Agric. Univ., 19(1):21-25.
- Sharma, L.C. and Sinha, S. (1974). Mycoflora of the rhizosphere of linseed in infected soil and its relation to root exudates. Proc. Nat. Acad. Sci. Indian, Sect. B., 44(3):176-185.
- Sheata, N.A., Afleger, F.L. and Davis, D.W. (1983). Response of susceptible and moderately resistant pea genotypes to interaction between Rhizoctonia solani and three other stem and root rot pathogens. Plant. Disease, 67(10):1146-1148.
- Shukla, D.S. and Misra, A.P. (1973). Effect of soil application of fungicides on inactivation of Sclerotia of Ozonium texanum var. parasiticum and incidence of gram wilt. Indian Phytopath., 25(3):378-386.



- Simmonds, P.M. and Codrington, R.J. (1937). A study of the fungus flora of wheat roots. Sci. Agric., 18(2):49-59.
- Singh, A.P. and Bhargava, S.N. (1981). Survival studies on the three species of Fusarium causing wilt of pigeonpea. (English summary). Phytopathologische Zeitschrift, 100(4):300-311.
- Singh, C. and Murty, Y.S. (1987). Studies on the effect of various growth regulators on seed germination and seedling growth of Cassia fistula L. Acta Bot. India, 15:206-212.
- ✓ Singh, D. (1971). Biology of soil fungi with emphasis on mycostasis. Ph.D. Thesis, Banaras Hindu University, Varanasi, India.
- Singh, D.B., Upadhyay, R.S. and Rai, B. (1978). Changes in amino acids and sugar contents of healthy and diseased levels of barley and roots of pigeonpea. Acta Bot. India, 6:154-158.
- Singh, G.N. (1981). Effect of foliar application of hormone on rhizosphere and rhizosphere and rhizoplane mycoflora of two oil yielding plants. Acta Bot. Indica, 9(2): 218-227.
- Singh, I. and Prasad, S.K. (1973). Effect of some nematicides on nematode and soil microorganisms. Indian Jour. Nematol., 3(2):109-133.
- Singh, N., Singh, R.P. and Lal, S. (1985). Effect of soil amendment on sugarcane wilt and its pathogens. Acta Bot. Indica, 13(2):212-217.
- Singh, P.N. (1970). Effect of foliar spray of hormones on rhizosphere mycoflora of Argemone maxicana Linn. Labdey Jour. Sci. Tech., SB:91-94.
- Singh, P.N., Sindhu, I.R. and Singhal, G. (1984). Fungi recorded from seeds and seedlings of Fagopyrum esculentum. Jour. Indian Bot. Soc., 63(3):236-243.
- Singh, U.P., Singh, N.B. and Singh, R.B. (1980). The fungicidal effect of neem (Azadirachta indica) extracts on some soil borne pathogens of gram Cicer arietinum. Mycologia, 72(6):1077-1093.

- Singh, N. and Singh, R.S. (1985). Inhibition of Fusarium udum (pigeonpea wilt pathogen) by ether distillate of Margosa cake amended soil. Indian Jour. Myco. and Plant Path., 13(3):329-330.
- Sinha, A.K. (1975). Control of Fusarium wilt of pigeonpea with bavistin, a systemic fungicide. Curr. Sci., 44(19): 700-701.
- Sinha, A.P., Agnihotri, V.P. and Singh, K. (1980). Effects of bavistin on soil microflora, their related physiological activity and growth of sugarbeet seedlings. Indian Jour. Exp. Biol., 18(5):489-491.
- Sinha, D.C. and Prasad, R.K. (1986). Effect of organic amendments in soil on foot and root rot of gram caused by Fusarium oxysporum f.sp. ciceri. Indian Phytopath., 39(1):145.
- Smith, N.R. (1948). Microbiology of soil. Ann. Rev. Microbiol., 2:435-484.
- Son, S.G., Shin, H.S. and Lee, M.W. (1985). Effect of amendments on ginseng root rot caused by Fusarium solani, population changes of microorganisms in soil. Korean Jour. Myco., 13(1):41-47.
- Sondhi, J.K. and Sinha, S. (1963). Rhizosphere mycoflora of Cicer arietinum L. Proc. 50th Indian Sci. Cong., p.347 (Abstr.).
- Srinivasan, K.V. (1968). The role of rhizosphere microflora in the resistant of sugarcane to Fusarium root rot. Proc. Int. Soc. Sugarcane, Technologists, 13th Congr., Taiwan, 2-17 March, 1968.
- Srivastava, L.S. and Dayal, R. (1981). Studies on rhizosphere mycoflora of Abelmoscus esculentus.X. Effect of fungicidal spray of rhizosphere and rhizoplane mycoflora. Indian Phytopath., 34(4):426-429.
- Srivastava, L.S. and Dayal, R. (1986). Studies on rhizosphere mycoflora of Abelmoscus esculentus XIV. Antibiosis by soil microorganisms. Indian Phytopath., 39(1):104-105.
- Srivastava, V.B. (1971). Investigation into rhizosphere microflora.VIII. Light and Dark treatments in relation to root region microflora. Plant and Soil, 35(3):463-470.

- ✓ Srivastava, V.B. (1974). Investigations into rhizosphere microflora IX. Effect of mixed cropping. Beitiefte Zur. Nova Hedwigia, 47:413-420.
- Srivastava, V.B. and Mishra, R.P. (1971). Investigation on the rhizosphere microflora II. Foliar application of some fungicides and antibiotics V. Effect of defoliation. Acta Phytopath. Acad. Sci. Hung., 5(2-4):303-308, 309-315.
- ✓ Srivastava, V.B. and Mishra, R.P. (1972). Rhizosphere fungal variation in susceptible and healthy wheat. Indian Phytopath., 24(4):784-786.
- Srivastava, V.B. and Sinha, S. (1971). Effect of various soil amendments on the wilt of coriander (Coriandrum sativum L.) Indian Jour. Agric. Sci., 41(9):779-782.
- Starkey, R.L. (1929a). Some influences on the development of higher plants upon the microorganisms in the soil.I. Historical and Introductory. Soil Sci., 27:319-334.
- Starkey, R.L. (1929b). Some influences on the development of higher plants upon the microorganisms in the soil.III. Influence of the stage of plant growth upon abundance of organisms. Soil Sci., 27:355-378.
- ✓ Starkey, R.L. (1929c). Some influences on higher plants upon the microorganisms in the soil.III. Influence of the stage of plant growth upon some activities of the organisms. Soil Sci., 27:433-444.
- Starkey, R.L. (1931). Some influences of higher plants upon the microorganisms in the soil.IV. Influence of the proximity to roots on abundance and activity of microorganisms. Soil Sci., 32:367-393.
- ✓ Starkey, R.L. (1958). Interrelations between microorganisms and plant roots in the rhizosphere. Bact. Rev., 22(3): 154-172.
- Stefurak, V.P. (1976). [Microflora in the rhizosphere of spruce stand infected by Fomitopsis annosa Karst.]. Mikrobiol.Z., 38(3):279-282.
- Stelfox, D. and Williams, J.R. (1980). Pythium species in alfalfa fields in central Alberta. Can. Plant Dis. Survey, 60(4):35-36.

- Stenton, H. (1958). Colonization of roots of Pisum sativum L. by fungi. Trans. Brit. Mycol. Soc., 41:74-80.
- Steward, F.C. and Durzan, D.J. (1965). Metabolism of nitrogenous compounds. In Plant Physiol. IVA (Edt. by Steward, F.C.), pp.379-679.
- <sup>+</sup>Strzeczkyk, E. (1961). Studies on the incidence of certain nutritional and physiological groups of bacteria in rhizosphere and non-rhizosphere soil. Acta Microbiol., Pol., 10:169-180.
- <sup>+</sup>Strzeczkyk, E. (1964). Studies on the rhizosphere microflora of plant resistant and susceptible to soil borne diseases II. Effect of amino acids and vitamins on growth of Thielaviopsis basicola (Berk. Br.) Ferr and Fusarium oxysporum f. lini (Bolley) Snyder et Hans. Acta Microbiol. Pol., 13(2):137-147.
- ✓ Subbarao, N.S. and Bailey, D.L. (1961). Rhizosphere studies in relation to varietal resistance of susceptibility of tomato to Verticillium wilt. Can. Jour. Bot., 39: 1747-1758.
- Sullia, S.B. (1968). Effect of foliar spray of hormones on the rhizosphere mycoflora of leguminous weeds. Plant and Soil, 29:292-298.
- Sullia, S.B. (1969). The fungicide kitazin and the mycoflora of Rice. Proc. Indian Acad. Sci. Sect. B., 69(6):295-308.
- ✓ Sullia, S.B. (1973). Effect of root exudates and extracts on rhizosphere fungi. Plant and Soil, 39(1):197-200.
- Sulochna, C.B. (1958). Root exudates. Mem. Indian Bot. So., 1:98-101.
- ✓ Tandon, A. and Tiwari, V.K. (1982). A comparative study of soil mycoflora under different crop covers. Acta Bot. Indica, 10:291-295.
- Tandon, R.N. (1970). Certain problems of post harvest diseases of fruits and vegetables. Indian Phytopath., 23:1-15.
- Thind, T.S. and Jhooty, J.S. (1985). Translocation and accumulation of benlate and bavistin in different parts of pea seedlings. Pesticides, 19(5):50-51.

- Thom, C. and Humfeld, H. (1932). Notes on the association of microorganisms and roots. Soil Sci., 34(1):29-36.
- Throne, G.N. (1954). Absorption of nitrogen, phosphorus, and potassium from nutrient sprays of leaves. Jour. Expt. Biol., 5:37-48.
- ✓ Timonin, N.I. (1940a). The interaction of higher plants and soil microorganisms.I. Microbiol population of rhizosphere of seedlings of certain cultivated plants. Can. Jour. Res. Sect. C., 18(7):307.
- Timonin, N.I. (1940b). The interaction of higher plants and soil microorganisms.II. Study of the microbial population of the rhizosphere in relation to resistance of plants to soil borne diseases. Can. Jour. Res. Sect. C., 18(9):444-455.
- Timonin, N.I. (1941). The interaction of higher plants and soil microorganisms.III. Effect of by-products of plant growth on activity of fungi and actinomycetes. Soil Sci., III., 5:395-408.
- ✓ Timonin, N.I. (1966). Rhizosphere effect of healthy and diseased lodgepole pine seedlings. Can. Jour. Microbiol., 12(3):531-537.
- Todorovic, M, Kalinovie and Tesic, Z. (1972). Investigation on the soil and rhizosphere microflora of maize in monoculture and crop rotation. Acta Biologica Iugoslavica B., 9(2):213-229.
- Tolle, R. and Rippel-Badles, A. (1958). [Studies on the rhizosphere of Gramineae]. Untersuchungen uber die Rhizosphere von Gramineen. Zbl. Bact. Abt., 2, 111(6-7): 204-217.
- Torstensson, L. and Wessen, B. (1984). Interaction between the fungicide benomyl and soil microorganisms. Soil Biol. and Biochem., 16(5):445-452.
- Tripathi, N.N. and Sharma, B.K. (1983). Incidence of chickpea dry root rot (Rhizoctonia bataticola) in Southern Haryana. Int. Chickpea Newsletter, 8:22-23.
- Tu, C.M. (1972). Effect of four nematicide on activities of microorganisms in soil. Appl. Microbiol., 23:398-401.

- Tu, C.M. (1973). The temperature dependent effect of residual nematicides on the activities of microorganisms. Can. Jour. Microbiol., 19(7):855-859.
- Upadhyay, R.S. and Rai, B. (1981). Effect of cultural practices and soil treatment on incidence of wilt disease of pigeonpea. Plant and Soil, 62(2):309-312.
- Upadhyay, R.S. and Rai, B. (1983). Competitive saprophytic ability of Fusarium udum in relation to some micro fungi of root region of pigeonpea. Indian Phytopath., 36(3): 539-544.
- Upadhyay, R.S. and Rai, B. (1986). Hyphal interaction and parasitism amongst some rhizosphere fungi of pigeonpea. Indian Phytopath., 36(2):302-306.
- Vaartaja, O. and Agnihotri, V.P. (1970). A comparison of rhizosphere flora in conifer beds treated with fungitoxicants. Zentbl. Bakt. Parasit. Kde Abt. 2, 124(2): 157-164.
- Vaidehi, B.K. (1973). Effect of foliar application of urea on the behaviour of Helminthosporium hawaiiensis on the rhizosphere of rice. Indian Jour. Myco. and Plant Path., 3(1):81-85.
- <sup>+</sup>Valaskova, Eva (1968). [The sensitivity of soil fungi to herbicides]. Die Empfindlichkeit Von Bodenpilzen gegenüber Herbiziden. Pflanzenschutzberichte, 38(10-11): 135-146.
- Vancura, V. and Hovadik, A. (1965). Root exudates of plants. II. Composition of root exudates of some vegetables. Plant and Soil, 22:21-32.
- Vardavakis, E. (1986). Vertical distribution and seasonal fluctuations of cellolytic fungi in a typic calcixeroll soil in Greece. Trans. Brit. Mycol. Soc., 86(4):668-672.
- Vargas, R. and Ramirez, C. (1983). [Biological control damping off caused by Rhizoctonia solani on cotton]. Combate biologico del Mal del talleulo causado per Rhizoctonia solani en algodón. Agronomia eastericense, 7(1/2): 73-75.
- Venkata Ram (1960). Foliar application of nutrients and rhizosphere microflora of Camellia sinensis Nature (Lond.), 187:621-622.

- Venkatesan, R. (1962). Studies on the actinomycetes population of paddy soil. Ph.D. Thesis, Dep. Agric. Annamalai Univ., Annamalaingor, South India.
- Venkatasubbaiah, P., Safeeulla, K.M. and Somashekar, R.K. (1984). Some metabolic changes in coffee seedlings due to Rhizoctonia solani infection. Life Sci. Adv., 3(2/3): 110-113.
- Venkatasubbaiah, P., Safeeulla, K.M. and Shetty, H.S. (1983). Phenolic content of coffee seedlings as influenced by Rhizoctonia solani. Indian Phytopath., 36(4):754-755.
- Verma, V. (1984). A text book of economic botany. p.62.
- Vesely, D. (1985). [Changes in the distribution of microfungi in the rhizosphere of sugarbeet]. Zmeny ve vyskytu Mikromycetu rhizosphere repy cukrove. Sbornik, UVTIZ Ochrana Roslin, 21(3):185-194.
- Vesely, D. (1986). [Relation between accompanying and plant pathogenic mycoflora of the rhizosphere of emerging sugarbeet]. Vzťahy medzi doprovodnou a fytopatogennou mycoflorou z rhizosphery Vzchzejici repy cukrove sbornik, UVTIZ Ochraňa Roslin, 22(2):109-118.
- Vir, D. and Sharma, R.K. (1985). Efficacy of fungicide:XXII. Evaluation of neem oil for control of pathogens. Asian Form Chemicals, 1(7/8):23-24.
- Vishwanath, N.R., Patel, R.B. and Rangaswami, G. (1969). Rhizosphere microflora of coffee plants affected by decline disease. Curr. Sci., 38(20):447-448.
- Vohra, S. and Chand, I.N. (1971). Relation of free amino acids to the development of angular leaf-spot (Blackarm) of cotton. Phytopath. Z., 70-177-180.
- Voronkevich, I.V., Matveeva, E.V. and Odintsova, M.A. (1972). The rhizosphere as the habitate of phytopathogenic bacteria.II. Significance of the rhizosphere in the life cycle of causal agents of soft rots. Biologicheskie Nauki, 15(9):103-109.
- Voroshilova, E.A. (1956). The effect of the apole free on the number of microorganisms. Microbiology (U.S.S.R.), 25:670-692 (English Translation).

- Vrany, J. (1963). Effect of foliar application of urea on the wheat microflora. Folia Microbiol., 8(6):351-355.
- Vrany, J. (1972). The effect of foliar application of urea on the root fungi of wheat growing in soil artificially contaminated with Fusarium spp. Foliar Microbiol., 17(6):500-504.
- Vrany, J., Vancura, V. and Macura, J. (1962). The effect of foliar application of some readily metabolised substances, growth regulators and antibiotics on rhizosphere microflora. Folia Microbiol., 7:61-70.
- ✓ Wadhwani, K. and Mehrotra, M. (1982). Fungi associated with roots of smut infected plants of Cynodon dactylon. Indian Phytopath., 35(4):722-724.
- Wadhwani, K. and Srivastava, M. (1985). The rhizosphere and rhizoplane fungi of coralloid roots of Cycas revoluta Thomb. Acta Bot. Indica, B (1):116-118.
- Wainwright, M. and Sowden, F.J. (1977). Influence of fungicide treatment on  $\text{CaCl}_2$  extractable phosphorus solubilizing microorganisms in soil. Plant and Soil, 48(2):335-345.
- Waksman, S.A. (1944). Three decades with soil fungi. Soil Sci., 58:89-115.
- Warcup, J.H. (1957). Studies on the occurrence and activity of fungi in wheat field soil. Trans. Brit. Myco. Soc., 40:237-262.
- Welte, E. and Trolldenier, G. (1961). Influence of the hydrogen ion concentration of soil on the rhizosphere effect. Naturewissenschaften, 48:509.
- West, P.M. (1939). Extraction of Thiamin and Biotin by roots of higher plants. Nature (Lond.), 144:1050-1051.
- + Winter, A.G. and Rumker, R.V. (1949). [The importance of the microflora of the rhizosphere in the resistance of roots to fungal diseases]. Die Bedeutung der microflora der wurzenaken zone fur die Resistenz von ivarzellen gegen. Pilzkhankhelten. Naturwiss., 36(1):30-31.
- enekowski, S. (1975). Studies on the influence of fungus communities of podzol soil on the growth of Fusarium oxysporum Schl. Prace. Naukowe Instytutu Ochrony, Roslin, 15(2):21-51.



- Wong, D.H., Sivasithampam, K. and Barbetti, M.J. (1986). Influence of soil temp., moisture and other fungal root pathogens on the pathogenicity of Phytophthora clandestina to subterranean clover. Trans. Brit. Myco. Soc., 86(3):479-482.
- Wood, F.W. (1960). Biological antagonism due to phytotoxic root exudates. Bot. Rev., 26:546-569.
- ✓ Wood, R.K.S. (1967). Physiological Plant Pathology. Blackwell Scientific Publications, Oxford and Edinburgh.
- Yarden, O. (1986). Fungicidal control of Sclerotinia sclerotiorum in soil with a combination of benomyl and thiram. Plant Disease, 70(8):738-742.
- Yasmeen, Prakash, D. and Khan, M.A.A. (1982). Rhizosphere and rhizoplane mycoflora of a medicinal plant. Jour. Sci. Res., A:37-40.
- Yih, Y. and Clark, H.C. (1965). Carbohydrate and protein contents of boron deficient tomato root tips. Plant Physiol., 40:312.
- +Youssef, B.A., Amr, A.M. and Heitefuss, R. (1985). [Interaction between herbicides and soil borne pathogens of cotton under greenhouse conditions] Wechselwirkungen Zwischen Herbiziden und bodenburtigen pathogenen Von Baumwolle unter Gewachs aubedingagen. Zeitschrift fur pflanzenkarkeiten und Pflanzenschutz, 92(1):55-63.
- Youssef, Y.A. and Mankarios, A.T. (1968). Studies on the rhizosphere mycoflora of broad bean and cotton.I. The fungal flora of the rhizosphere. Mycopath. Myco. Appl., 35:389-400.
- Youssef, Y.A. and Mankarios, A.T. (1969). Studies on the rhizosphere mycoflora of broad bean and cotton.II. Seed and root exudates and their effects on spore germination and growth of prevalent fungi isolated from the rhizosphere. Mycopath. Mycol. Appl., 38(3): 257-269.
- Youssef, Y.A. and Mankarios, A.T. (1974). Studies on rhizosphere mycoflora of broad bean and cotton.IV. The influence of the rhizosphere fungi on plant grwoth. Mycopath. Mycol. Appl., 54(2):173-180.

- Yuen, G.Y. and Schroth, M.N. (1986). Interactions of Pseudomonas fluorescens strains E6 with ornamental plants and its effect on the composition of root colonizing microflora. Phytopath., 76(1):176-180.
- Zagallo, A.C. and Bollen, W.B. (1962). Studies on the rhizosphere of tall fescus. Ecology, 43:54-62.
- Zakaria, M.A., Lockwood, J.L. and Filonow, A.B. (1980). Reduction in Fusarium population density in soil by volatile degradation products of oilseed meal amendments. Phytopath., 70(6):495-499.
- Zgorkiewicz, A. and Mackiewicz, S. (1975). Trials of the elaborated method for preventing the appearance of Fusarium diseases of Lupin involving the use of different crop rotations on different types of soil. Proc. Naukowe Instutu Ochrany Roslin, 17(2):61-95.
- <sup>+</sup>Zhdanova, N.N. (1963). The occurrence of dark hyphomycete in the rhizosphere of maize in steppes and forest steppes of ukraine. Mycrobiol., Zh., 25(4):2834.
- <sup>+</sup>Zhukovskaya, S.A. (1976). Role of soil fungi in the development of diseases of soybean in the Primorsk region. Tr. Biot. Bochr. Inst. Dalnevost. Nauch. Tsentre ANSSSR, 41:66-83.
- Zombolin, L. and Schenck, N.C. (1983). Reduction of the effect of pathogenic, root infecting fungi on soybean by the mycorrhizal fungus Glomus mossca. Phytopath., 73(10):1403.
- Zozzerini, A. and Tosi, L. (1985). Antagonistic activity of fungi isolated from Sclerotia of Sclerotinia sclerotiorum Plant Path., 34(3):415-421.

---

<sup>+</sup>Original not seen